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(54) Title: VIRAL ENCODED SEMAPHORIN PROTEIN RECEPTOR DNA AND POLYPEPTIDES			
(57) Abstract The invention is directed to VESPR polypeptides as a purified and isolated protein, the DNA encoding the VESPR polypeptide, host cells transfected with cDNAs encoding VESPR, and methods for preparing VESPR polypeptides.			

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**VIRAL ENCODED SEMAPHORIN PROTEIN RECEPTOR DNA AND
POLYPEPTIDES**

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FIELD OF THE INVENTION

The present invention relates to semaphorin receptor polypeptides, the nucleic acids encoding such semaphorin receptor polypeptides, processes for producing recombinant semaphorin receptor polypeptides, and pharmaceutical compositions containing such polypeptides.

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BACKGROUND OF THE INVENTION

The semaphorin gene family includes a large number of molecules that encode related transmembrane and secreted glycoproteins known to be neurologic regulators. The semaphorins are generally well conserved in their extracellular domains which are typically about 500 amino acids in length. Semaphorin family proteins have been observed in neuronal and nonneuronal tissue and have been studied largely for their role in neuronal growth cone guidance. For example, the secreted semaphorins known as collapsin-1 and Drosophila semaphorin II are selectively involved in repulsive growth cone guidance during development. Flies having semaphorin II genes that are mutated so that their function is reduced exhibit abnormal behavior characteristics.

Another semaphorin gene has been identified in several strains of poxvirus. This semaphorin is found in vaccinia virus (Copenhagen strain) and is encoded in an open reading frame (ORF) known as A39R. The A39R encoded protein has no transmembrane domain and no potential membrane linkage and is known to be a secreted protein. A variola virus ORF also contains sequences that share homology with the vaccinia virus ORF A39R at the nucleotide level and the amino acid level. Another viral semaphorin, AHV-sema, has been found in the Alcelaphine Herpesvirus (AHV).

Genes encoding mammalian (human, rat, and mouse) semaphorins have been identified, based upon their similarity to insect semaphorins. Functional studies of these semaphorins suggest that embryonic and adult neurons require a semaphorin to establish workable connections. Significantly, the fast response time of growth cone cultures to appropriate semaphorins suggests that semaphorin signaling involves a receptor-mediated signal transduction mechanism. To date, one semaphorin receptor, designated neuropilin, has been isolated using mRNA from rat spinal cord. Another

5 receptor, designated neuropilin-2, has been suggested (Kolodkin et al. *Cell* 90:753-762, 1997)

Semaphorin ligands that are secreted into the extracellular milieu signal through receptor bearing cells in a local and systemic fashion. In order to further investigate the nature of cellular processes regulated by such local and systemic
10 signaling, it would be beneficial to identify additional semaphorin receptors and ligands. Furthermore, because virus encoded semaphorins are produced by infected cells and are present in viruses that are lytic (poxviruses) and viruses that are not known to be neurotropic (AHV), it is unlikely that their primary function is to modify neurologic responses. It is more likely that the virus encoded semaphorins function to
15 modify the immunologic response of the infected host and it is likely that mammalian homologues to virus encoded semaphorins function to modify the immunologic response. In view of the suggestion that viral semaphorins may function in the immune system as natural immunoregulators it would be beneficial to identify semaphorin receptors as therapeutic agents for enhancing or downregulating the
20 immune response.

SUMMARY OF THE INVENTION

The present invention pertains to semaphorin receptors as isolated or homogeneous proteins. In particular, the present invention provides a semaphorin
25 receptor polypeptide, designated VESPR (Viral Encoded Semaphorin Protein Receptor) that binds semaphorins, including, but not limited to, the A39R vaccinia semaphorin and AHV semaphorin. Also, within the scope of the present invention are DNAs encoding VESPR polypeptides and expression vectors that include DNA encoding VESPR polypeptides. The present invention also includes host cells that
30 have been transfected or transformed with expression vectors that include DNA encoding a VESPR polypeptide, and processes for producing VESPR polypeptides by culturing such host cells under conditions conducive to expression. The present invention further includes antibodies directed against VESPR polypeptides.

Further within the scope of the present invention are processes for purifying or
35 separating semaphorins or cells that express semaphorins to which the VESPR polypeptides of the present invention bind. Such processes include binding at least one VESPR polypeptide to a solid phase matrix and contacting a mixture containing a semaphorin polypeptide to which the VESPR polypeptide binds, or a mixture of cells expressing the semaphorin with the bound VESPR polypeptide, and then separating
40 the contacting surface and the solution.

5 The present invention additionally provides processes for treating inflammation and inflammatory diseases. Such processes include administering a therapeutically effective amount of a soluble VESPR polypeptide to an human or other mammal afflicted with a disease associated with proinflammatory activity of a semaphorin ligand.

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DETAILED DESCRIPTION OF THE INVENTION

The present invention provides novel semaphorin receptor polypeptides designated Viral Encoded Semaphorin Protein Receptor (VESPR), DNA encoding
15 VESPR polypeptides and recombinant expression vectors that include DNA encoding VESPR polypeptides. The present invention additionally provides methods for isolating VESPR polypeptides and methods for producing recombinant VESPR polypeptides by cultivating host cells transfected with the recombinant expression vectors under conditions appropriate for expressing semaphorin receptors and
20 recovering the expressed receptor polypeptide.

In particular, the present invention provides VESPR polypeptides that bind semaphorins, including but not limited to, the vaccinia virus A39R semaphorin and the AHV semaphorin. The native VESPR polypeptide described herein was isolated using an Ectromelia virus A39R semaphorin/Fc fusion protein (A39R/Fc) to recover
25 VESPR from the membranes of human cells expressing the receptor. As described in the examples below, flow cytometry experiments establish that the VESPR polypeptide polypeptides of the present invention are expressed by B cells lines, monocyte-type cell lines, T cell lines, dendritic cells NK cells, lung epithelial cells, stroma, intestinal epithelial cells and lymphoma cells.

30 Furthermore, as demonstrated in the examples below, VESPR polypeptides of the present invention bind with their ligands to participate in upregulating the CD69 activation antigen on dendritic cells. Also characteristic of semaphorin receptors described herein is their ability to interact with their ligands to synergize with interferon and SAC to upregulate IL-12 production and down regulate MHC class II
35 and CD86 expression on mouse dendritic cells. VESPR polypeptides of the present invention are also associated with increased expression of CD54 on monocytes which suggests cellular activation as a result of the interaction between semaphorins and their receptors. Among the uses of the VESPR polypeptides that flow from aforementioned biological properties of the receptor-ligand interaction are inducing
40 IL-12 production and subsequent natural killer cell activation. VESPR polypeptides find further use in treating diseases and adverse conditions associated with

5 inflammation. In particular, soluble VESPR polypeptides can be used to antagonize proinflammatory activities associated with the interaction of semaphorin ligands and their receptors. Rheumatoid arthritis, a disease associated with chronic inflammation of synovial tissue, has been linked with upregulation of the human semaphorin E gene (Mangasser-Stephan et al., *Biochem and Biophys Res Com*, 234:153-156, 1997).
10 Thus, soluble forms of VESPR polypeptides of the present invention may be useful in downregulating semaphorin activity that mediates this inflammatory disease.

VESPR, a native semaphorin receptor of the present invention, was isolated using a viral semaphorin ligand known as Ectromelia A39R. Example 1 below describes isolating the A39R semaphorin ligand and preparing an A39R/Fc fusion
15 protein which was used to identify cell lines that bind the ligand and to determine the effects of interactions between A39R and its cell bound receptor.

Examples 4 and 5 describe identifying a native VESPR polypeptide of the present invention and isolating and purifying a human VESPR polypeptide. The amino acid sequence of the human VESPR polypeptide, isolated as described in
20 Example 5, is disclosed in SEQ ID NO:2. The amino acid sequence of SEQ ID NO: 2 was obtained by sequencing the isolated and purified receptor using tandem mass spectrometry analysis of peptides produced in a trypsin digestion, in combination with contiguous EST sequences and identified cDNAs. The amino acid sequence presented in SEQ ID NO:2 has a predicted extracellular domain of amino acids 1-944
25 that includes a signal peptide with a cleavage site predicted at amino acid 34. The predicted transmembrane domain of SEQ ID NO:2 includes amino acids 945-965 and the cytoplasmic domain of SEQ ID NO:2 extends from amino acids 966-1568.

A DNA encoding amino acids 19-1100 of human VESPR in *E. coli* DH10B was deposited with the American type Culture Collection, Rockville, MD, USA on
30 _____ and assigned accession number _____. The deposit was made under the terms of the Budapest Treaty. The DNA construct of the deposit differs from that of SEQ ID NO: 1 in that nucleotide 172 is C. The resulting encoded amino acid 58 is leu.

Amino acid sequence searches were performed in available data bases for
35 proteins and polypeptides sharing homology with the full length VESPR or domains thereof. The searches for polypeptides sharing homology with VESPR were performed using the BLAST algorithm described by Altschul et al., *J Mol Bio* 215:403-410 (1990). This program was used to compare the VESPR amino acid sequence with protein and DNA sequences found in data bases obtained from the
40 National Center for Biotechnology Information. Similarity scores obtained as a result

5 of these searches identified groups of polypeptides having varying degrees of homology with VESPR. The highest degree of similarity was found to be between the VESPR and a group of proteins known as the "plexin gene family" (Maestrini et al., 1996, and Kameyama et al., 1996). Pairwise and multiple sequence alignments between VESPR and human and murine members of the plexin family were
10 performed using the Smith-Waterman algorithm as implemented in the Genetics Computer Group (GCG) programs "BESTFIT" and "PILEUP" (Wisconsin Package, 9.0). The GCG program "DISTANCES" was used to calculate average pairwise percentage identity of the aligned protein sequences.

Pairwise sequence alignments between the VESPR polypeptide and each of
15 several members of the plexin gene family revealed an average identity in their cytoplasmic domain (amino acids 966-1568) of 39% to 40% and an average identity for each of the entire protein of 24% - 25%. The higher degree of homology in the cytoplasmic domains suggests similar signal transduction mechanisms among the cytoplasmic domains.

20 In order to identify regions of similarity between the protein sequences found to have some overall homology, homology analyses of the results of protein data base searches were performed using the BLIXEM and MSPCRUNCH programs (Sonnhammer and Durbin (1944a,b). The homology analyses revealed a novel subdomain with similarity to a region of the semaphorin domain of a number of
25 members of the semaphorin family of genes described by Kolodkin et al. (1993). The novel subdomain includes amino acids 380-482 of the VESPR sequence of the present invention. This subdomain can be subdivided into two distinct smaller regions, that include residues 388-402 and 454-482, respectively. The C-proximal half-subdomain contains several highly conserved cysteine and tryptophan residues,
30 forming a consensus sequence of C-x(5)-C-x(2)-C-x(7)-C-x-W-C-x(5)-C, where x is any amino acid. This entire subdomain is distinct from the canonical semaphorin domain described for the semaphorin gene family in that (a) it is smaller (100 amino acid residues for the subdomain vs 500 residues for the entire semaphorin domain), (b) it is also present in the plexin gene family and MET-hepatocyte growth factor
35 receptor family, neither of which is a canonical semaphorin gene family members, and (c). it is present in a gene which is not itself a member of the semaphorin gene family but which interacts with a member of the semaphorin family (A39R). These subdomain sequences, therefore, represent peptides that are potentially capable of further identifying other receptors which interact with semaphorins.

5 A cDNA sequence that encodes the VESPR polypeptide of SEQ ID NO:2 was assembled as a composite of contiguous EST and cloned cDNA nucleotide sequences and is disclosed in SEQ ID NO:1. As described in Example 5, identifying the cDNA that encodes the amino acid sequence of SEQ ID NO:2 enables constructing expression vectors that include the encoding cDNAs. Then culturing host cells
10 transfected with a recombinant expression vector that contains cDNA encoding VESPR polypeptide, under conditions appropriate for expressing the VESPR polypeptide, and recovering the expressed VESPR polypeptide provides methods for producing VESPR polypeptides of the present invention.

15 Since VESPR polypeptide is found in B cell lines, T cell lines and dendritic cells, treating a variety of conditions associated with overactive or underactive immuno-regulation is possible. Moreover, the ligand and receptor complex may be involved in neural growth, development and/or maintenance. While not limited to such, particular uses of the VESPR are described infra.

20 The terms "VESPR" and "VESPR polypeptide" of the present invention encompass polypeptides having the amino acid sequence SEQ ID NO:2, and proteins that are encoded by nucleic acids that contain the nucleic acid sequence of SEQ ID NO:1. In addition, the terms include those polypeptides that have a high degree of similarity or a high degree of identity with the amino acid sequence of SEQ ID NO:2, which polypeptides are biologically active and bind at least one molecule or
25 fragments of a molecule that are members of the semaphorin family. In addition, the term VESPR refers to biologically active gene products of the DNA of SEQ ID NO:1. Further encompassed by the term VESPR are soluble or truncated proteins that comprise primarily the binding portion of the protein, retain biological activity and are capable of being secreted. Specific examples of such soluble proteins are those
30 comprising the sequence of amino acids 1-944 of SEQ ID NO:2.

The term "biologically active" as it refers to VESPR or semaphorin receptor polypeptide, means that the VESPR or semaphorin receptor polypeptide is capable of binding to at least one semaphorin. Assays suitable for determining VESPR binding are described herein and can include standard flow cytometry tests and slide binding
35 tests.

"Isolated" means a VESPR is substantially free of association with other proteins or polypeptides residual of the expression process, for example, as a purification product of recombinant host cell culture or as a purified extract.

40 A VESPR variant as referred to herein, means a polypeptide substantially homologous to native VESPR, but which has an amino acid sequence different from

5 that of native VESPR because of one or more deletions, insertions or substitutions. The variant amino acid sequence preferably is at least 80% identical to a native VESPR amino acid sequence, most preferably at least 90% identical. The percent identity may be determined, for example, by comparing sequence information using the GAP computer program, version 8.1 described by Devereux et al. (*Nucl. Acids*
10 *Res.* 12:387, 1984) and available from the University of Wisconsin Genetics Computer Group (UWGCG). The preferred default parameters for the GAP program include: (1) a unary comparison matrix (containing a value of 1 for identities and 0 for non-identities) for nucleotides, and the weighted comparison matrix of Gribskov and Burgess, *Nucl. Acids Res.* 14:6745, 1986, as described by Schwartz and Dayhoff,
15 eds., *Atlas of Protein Sequence and Structure*, National Biomedical Research Foundation, pp. 353-358, 1979; (2) a penalty of 3.0 for each gap and an additional 0.10 penalty for each symbol in each gap; and (3) no penalty for end gaps. Variants may comprise conservatively substituted sequences, meaning that a given amino acid residue is replaced by a residue having similar physiochemical characteristics.
20 Examples of conservative substitutions include substitution of one aliphatic residue for another, such as Ile, Val, Leu, or Ala for one another, or substitutions of one polar residue for another, such as between Lys and Arg; Glu and Asp; or Gln and Asn. Other such conservative substitutions, for example, substitutions of entire regions having similar hydrophobicity characteristics, are well known. Naturally occurring
25 VESPR variants or alleles are also encompassed by the invention. Examples of such variants are proteins that result from alternate mRNA splicing events or from proteolytic cleavage of the VESPR protein, wherein the binding property is retained. Alternate splicing of mRNA may yield a truncated but biologically active VESPR polypeptide, such as a naturally occurring soluble form of the protein, for example.
30 Variations attributable to proteolysis include, for example, differences in the N-termini upon expression in different types of host cells, due to proteolytic removal of one or more terminal amino acids from the VESPR polypeptide (generally from 1-5 terminal amino acids).

As mentioned above, Example 1 describes the construction of novel viral
35 A39R/Fc fusion proteins useful in studying VESPR binding. Other antibody Fc regions may be substituted for the human IgG1 Fc region described in the Example. Suitable Fc regions are those that can bind with high affinity to protein A or protein G, and include the Fc region of human IgG1 or fragments of the human or murine IgG1 Fc region, e.g., fragments comprising at least the hinge region so that interchain
40 disulfide bonds will form. The viral A39R:Fc fusion protein offers the advantage of

5 being easily purified. In addition, disulfide bonds form between the Fc regions of two separate fusion protein chains, creating dimers.

As described above, in one aspect, the present invention includes soluble VESPR polypeptides. Soluble VESPR polypeptides comprise all or part of the extracellular domain of a native VESPR but lack the transmembrane region that
10 would cause retention of the polypeptide on a cell membrane. Soluble VESPR polypeptides advantageously comprise the native (or a heterologous) signal peptide when initially synthesized to promote secretion, but the signal peptide is cleaved upon secretion of VESPR polypeptides from the cell. Soluble VESPR polypeptides encompassed by the invention retain the ability to bind semaphorin ligands. Indeed,
15 soluble VESPR polypeptides may also include part of the signal or part of the cytoplasmic domain or other sequences, provided that the soluble VESPR protein can be secreted.

Soluble VESPR may be identified (and distinguished from its non-soluble membrane-bound counterparts) by separating intact cells which express the desired
20 protein from the culture medium, e.g., by centrifugation, and assaying the medium (supernatant) for the presence of the desired protein. The presence of VESPR in the medium indicates that the protein was secreted from the cells and thus is a soluble form of the desired protein.

Soluble forms of VESPR polypeptides possess many advantages over the
25 native, membrane bound VESPR protein. Purification of the proteins from recombinant host cells is feasible, since the soluble proteins are secreted from the cells. Further, soluble proteins are generally more suitable for intravenous administration.

Examples of soluble VESPR polypeptides include those comprising a
30 substantial portion of the extracellular domain of a native VESPR polypeptide. An example of a soluble VESPR polypeptide is amino acids 1-944 of SEQ ID NO:2. In addition, truncated soluble VESPR proteins comprising less than the entire extracellular domain are included in the invention, e.g. amino acids 35-944. When initially expressed within a host cell, soluble VESPR polypeptides may additionally
35 comprise one of the heterologous signal peptides described below that is functional within the host cells employed. Alternatively, the protein may comprise the native signal peptide. In one embodiment of the invention, soluble VESPR can be expressed as a fusion protein comprising (from N- to C-terminus) the yeast α -factor signal peptide, a FLAG[®] peptide described below and in U.S. Patent No. 5,011,912, and
40 soluble VESPR polypeptide consisting of amino acids 1-944 or 35-944 of SEQ ID

5 NO:2. This recombinant fusion protein is expressed in and secreted from yeast cells. The FLAG[®] peptide facilitates purification of the protein, and subsequently may be cleaved from the soluble VESPR using bovine mucosal enterokinase. Isolated DNA sequences encoding soluble VESPR proteins are encompassed by the invention.

10 Truncated VESPR polypeptides, including soluble polypeptides, may be prepared by any of a number of conventional techniques. A desired DNA sequence may be chemically synthesized using techniques known per se. DNA fragments also may be produced by restriction endonuclease digestion of a full length cloned DNA sequence, and isolated by electrophoresis on agarose gels. Linkers containing
15 restriction endonuclease cleavage site(s) may be employed to insert the desired DNA fragment into an expression vector, or the fragment may be digested at cleavage sites naturally present therein. The well known polymerase chain reaction procedure also may be employed to amplify a DNA sequence encoding a desired protein fragment. As a further alternative, known mutagenesis techniques may be employed to insert a stop codon at a desired point, e.g., immediately downstream of the codon for the last
20 amino acid of the binding domain.

As stated above, the invention provides isolated or homogeneous VESPR polypeptides, both recombinant and non-recombinant. Variants and derivatives of native VESPR proteins that retain the desired biological activity (e.g., the ability to bind to semaphorins) may be obtained by mutations of nucleotide sequences coding
25 for native VESPR polypeptides. Alterations of the native amino acid sequence may be accomplished by any of a number of conventional methods. Mutations can be introduced at particular loci by synthesizing oligonucleotides containing a mutant sequence, flanked by restriction sites enabling ligation to fragments of the native sequence. Following ligation, the resulting reconstructed sequence encodes an analog
30 having the desired amino acid insertion, substitution, or deletion.

Alternatively, oligonucleotide-directed site-specific mutagenesis procedures can be employed to provide an altered gene wherein predetermined codons can be altered by substitution, deletion or insertion. Exemplary methods of making the alterations set forth above are disclosed by Walder et al. (*Gene* 42:133, 1986); Bauer et al. (*Gene* 37:73, 1985); Craik (*BioTechniques*, January 1985, 12-19); Smith et al. (*Genetic Engineering: Principles and Methods*, Plenum Press, 1981); Kunkel (*Proc. Natl. Acad. Sci. USA* 82:488, 1985); Kunkel et al. (*Methods in Enzymol.* 154:367, 1987); and U.S. Patent Nos. 4,518,584 and 4,737,462 all of which are incorporated by
35 reference.

5 Native VESPR polypeptide may be modified to create VESPR derivatives by forming covalent or aggregative conjugates with other chemical moieties, such as glycosyl groups, lipids, phosphate, acetyl groups and the like. Covalent derivatives of VESPR polypeptides may be prepared by linking the chemical moieties to functional groups on VESPR amino acid side chains or at the N-terminus or C-terminus of a VESPR polypeptide or the extracellular domain thereof. Other derivatives of VESPR polypeptides within the scope of this invention include covalent or aggregative conjugates of VESPR polypeptides or its fragments with other proteins or polypeptides, such as by synthesis in recombinant culture as N-terminal or C-terminal fusions. For example, the conjugate may comprise a signal or leader polypeptide sequence (e.g. the α -factor leader of *Saccharomyces*) at the N-terminus of a VESPR polypeptide. The signal or leader peptide co-translationally or post-translationally directs transfer of the conjugate from its site of synthesis to a site inside or outside of the cell membrane or cell wall.

20 VESPR polypeptide fusions can comprise peptides added to facilitate purification and identification of VESPR. Such peptides include, for example, poly-His or the antigenic identification peptides described in U.S. Patent No. 5,011,912 and in Hopp et al., *Bio/Technology* 6:1204, 1988.

25 The invention further includes VESPR with or without associated native-pattern glycosylation. VESPR polypeptide expressed in yeast or mammalian expression systems (e.g., COS-7 cells) may be similar to or significantly different from a native VESPR polypeptide in molecular weight and glycosylation pattern, depending upon the choice of expression system. Expression of VESPR polypeptides in bacterial expression systems, such as *E. coli*, provides non-glycosylated molecules.

30 Equivalent DNA constructs that encode various additions or substitutions of amino acid residues or sequences, or deletions of terminal or internal residues or sequences not needed for biological activity or binding are encompassed by the invention. For example, N-glycosylation sites in the VESPR extracellular domain can be modified to preclude glycosylation, allowing expression of a reduced carbohydrate analog in mammalian and yeast expression systems. N-glycosylation sites in eukaryotic polypeptides are characterized by an amino acid triplet Asn-X-Y, wherein X is any amino acid except Pro and Y is Ser or Thr. The native human VESPR protein comprises 24 such triplets, at amino acids 86-88, 141-143, 149-151, 241-243, 252-254, 386-388, 407-409, 548-550, 553-555, 582-584, 588-590, 591-593, 653,655, 686-688, 692-694, 715-717, 741-743, 771-773, 796-798, 821-823, 871-873, 890-892, 895-897 and 920-922 of SEQ ID NO:2. Appropriate substitutions, additions or

40

5 deletions to the nucleotide sequence encoding these triplets will result in prevention of attachment of carbohydrate residues at the Asn side chain. Alteration of a single nucleotide, chosen so that Asn is replaced by a different amino acid, for example, is sufficient to inactivate an N-glycosylation site. Known procedures for inactivating N-glycosylation sites in proteins include those described in U.S. Patent 5,071,972 and
10 EP 276,846, hereby incorporated by reference.

In another example, sequences encoding Cys residues that are not essential for biological activity can be altered to cause the Cys residues to be deleted or replaced with other amino acids, preventing formation of incorrect intramolecular disulfide bridges upon renaturation. Other equivalents are prepared by modification of
15 adjacent dibasic amino acid residues to enhance expression in yeast systems in which KEX2 protease activity is present. EP 212,914 discloses the use of site-specific mutagenesis to inactivate KEX2 protease processing sites in a protein. KEX2 protease processing sites are inactivated by deleting, adding or substituting residues to alter Arg-Arg, Arg-Lys, and Lys-Arg pairs to eliminate the occurrence of these
20 adjacent basic residues. Lys-Lys pairings are considerably less susceptible to KEX2 cleavage, and conversion of Arg-Lys or Lys-Arg to Lys-Lys represents a conservative and preferred approach to inactivating KEX2 sites. The human VESPR contains 11 KEX2 protease processing sites

Nucleic acid sequences within the scope of the invention include isolated
25 DNA and RNA sequences that hybridize to the VESPR nucleotide sequences disclosed herein under conditions of moderate or high stringency, and that encode biologically active VESPR. Conditions of moderate stringency, as defined by Sambrook et al. *Molecular Cloning: A Laboratory Manual*, 2 ed. Vol. 1, pp. 101-104, Cold Spring Harbor Laboratory Press, (1989), include use of a prewashing
30 solution of 5 X SSC, 0.5% SDS, 1.0 mM EDTA (pH 8.0) and hybridization conditions of about 55°C, 5 X SSC, overnight. Conditions of high stringency include higher temperatures of hybridization and washing. The skilled artisan will recognize that the temperature and wash solution salt concentration may be adjusted as necessary according to factors such as the length of the nucleic acid molecule and the
35 relative amount of A, T/U, C and G nucleotides.

Due to the known degeneracy of the genetic code wherein more than one codon can encode the same amino acid, a DNA sequence may vary from that shown in SEQ ID NO:1 and still encode a VESPR polypeptide having the amino acid sequence of SEQ ID NO:2. Such variant DNA sequences may result from silent

5 mutations (e.g., occurring during PCR amplification), or may be the product of deliberate mutagenesis of a native sequence.

The invention provides equivalent isolated DNA sequences encoding biologically active VESPR, selected from: (a) cDNA comprising the nucleotide sequence presented in SEQ ID NO:1; (b) DNA capable of hybridization to a DNA of
10 (a) under moderately stringent conditions and that encodes biologically active VESPR polypeptide; (c) DNA that is degenerate as a result of the genetic code to a DNA defined in (a) or (b) and that encodes biologically active VESPR polypeptide; and (d) DNA complementary to the DNA of (a), (b) or (c). VESPR polypeptides encoded by such DNA equivalent sequences are encompassed by the invention.

15 DNAs that are equivalents to the DNA sequence of SEQ ID NO:1 will hybridize under moderately and highly stringent conditions to the DNA sequence that encodes polypeptides comprising the sequence of SEQ ID NO:2. Examples of VESPR proteins encoded by such DNA, include, but are not limited to, VESPR fragments and VESPR proteins comprising inactivated N-glycosylation site(s),
20 inactivated KEX2 protease processing site(s), or conservative amino acid substitution(s), as described above. VESPR polypeptides encoded by DNA derived from other species, wherein the DNA will hybridize to the cDNA of SEQ ID NO:1 are also encompassed.

Variants possessing the requisite ability to bind semaphorins may be identified
25 by any suitable assay. Biological activity of VESPR polypeptides may be determined, for example, by competition for binding to the receptor binding domain of semaphorins (i.e. competitive binding assays).

One type of a competitive binding assay for a VESPR polypeptide uses a radiolabeled, soluble VESPR and intact semaphorin-expressing cells. Instead of
30 intact cells, one could substitute soluble semaphorin:Fc fusion proteins bound to a solid phase through the interaction of a Protein A, Protein G or an antibody to the semaphorin or Fc portions of the molecule, with the Fc region of the fusion protein. Another type of competitive binding assay utilizes radiolabeled soluble semaphorins such as a fusion protein, and intact cells expressing VESPR.

35 Competitive binding assays can be performed following conventional methodology. In one embodiment, a soluble VESPR polypeptide can be made to compete with an immobilized receptor for binding with a soluble semaphorin ligand. For example, a radiolabeled soluble semaphorin ligand can be antagonized by soluble VESPR in an assay for binding activity against a surface-bound semaphorin receptor.

5 Qualitative results can be obtained by competitive autoradiographic plate binding assays, or Scatchard plots may be utilized to generate quantitative results.

Alternatively, semaphorin binding proteins, such as VESPR or anti-semaphorin antibodies, can be bound to a solid phase such as a column chromatography matrix or a similar substrate suitable for identifying, separating or
10 purifying cells that express semaphorin on their surface. Binding of a semaphorin-binding protein to a solid phase contacting surface can be accomplished by any means, for example, by constructing a VESPR:Fc fusion protein and binding such to the solid phase through the interaction of Protein A or Protein G. Various other means for fixing proteins to a solid phase are well known in the art and are suitable
15 for use in the present invention. For example, magnetic microspheres can be coated with VESPR and held in the incubation vessel through a magnetic field. Suspensions of cell mixtures containing semaphorin-expressing cells are contacted with the solid phase that has VESPR polypeptides thereon. Cells having semaphorin on their surface bind to the fixed VESPR and unbound cells then are washed away. This
20 affinity-binding method is useful for purifying, screening or separating such semaphorin-expressing cells from solution. Methods of releasing positively selected cells from the solid phase are known in the art and encompass, for example, the use of enzymes. Such enzymes are preferably non-toxic and non-injurious to the cells and are preferably directed to cleaving the cell-surface binding partner. In the case of
25 semaphorin-VESPR interactions, the enzyme preferably would cleave the semaphorin, thereby freeing the resulting cell suspension from the "foreign" semaphorin receptor material. The purified cell population then may be used to repopulate mature (adult) tissues.

Alternatively, mixtures of cells suspected of containing semaphorin-positive
30 cells first can be incubated with biotinylated VESPR. Incubation periods are typically at least one hour in duration to ensure sufficient binding to semaphorin. The resulting mixture then is passed through a column packed with avidin-coated beads, whereby the high affinity of biotin for avidin provides the binding of the cell to the beads. Use of avidin-coated beads is known in the art. See Berenson, et al. *J. Cell. Biochem.*,
35 10D:239 (1986). Washing unbound material and releasing the bound cells is performed using conventional methods.

As described above, VESPR can be used to separate cells expressing semaphorin. In an alternative method, VESPR or an extracellular domain or a fragment thereof can be conjugated to a detectable moiety such as ^{125}I to detect
40 semaphorin-expressing cells. Radiolabeling with ^{125}I can be performed by any of

5 several standard methodologies that yield a functional ^{125}I -VESPR molecule labeled to high specific activity. Or an iodinated or biotinylated antibody against the semaphorin receptor can be used. Another detectable moiety such as an enzyme that can catalyze a colorimetric or fluorometric reaction, biotin or avidin may be used. Cells to be tested for semaphorin-expression can be contacted with labeled VESPR
10 polypeptide. After incubation, unbound labeled VESPR is removed and binding is measured using the detectable moiety.

The binding characteristics of VESPR (including variants) may also be determined using a conjugated semaphorin (for example, ^{125}I -semaphorin:Fc) in competition assays similar to those described above. In this case, however, intact
15 cells expressing semaphorins bound to a solid substrate are used to measure the extent to which a sample containing a putative VESPR variant competes for binding with a conjugated semaphorin.

Other means of assaying for VESPR include the use of anti-VESPR antibodies, cell lines that proliferate in response to VESPR, or recombinant cell lines
20 that express semaphorin and proliferate in the presence of VESPR.

The VESPR proteins disclosed herein also may be employed to measure the biological activity of semaphorin proteins in terms of their binding affinity for VESPR. As one example, VESPR polypeptides of the present invention may be used in determining whether biological activity is retained after modification of a
25 semaphorin protein (e.g., chemical modification, truncation, mutation, etc.). The biological activity of a semaphorin protein thus can be ascertained before it is used in a research study, or in the clinic, for example.

VESPR polypeptides of the present invention find use as reagents that may be employed by those conducting "quality assurance" studies, e.g., to monitor shelf life
30 and stability of semaphorin protein under different conditions. To illustrate, VESPR polypeptides may be employed in a binding affinity study to measure the biological activity of an semaphorin protein that has been stored at different temperatures, or produced in different cell types. The binding affinity of the modified semaphorin protein for VESPR is compared to that of an unmodified semaphorin protein to detect
35 any adverse impact of the modifications on biological activity of the semaphorin.

VESPR polypeptides also find use as carriers for delivering agents attached thereto to cells expressing semaphorins. As described in example 7 below, a putative human semaphorin is expressed in cells found in the placenta, testis, ovary and spleen. VESPR polypeptides can thus can be used to deliver diagnostic or therapeutic

5 agents to these cells (or to other cell types found to express a semaphorin on a cell surfaces) in *in vitro* or *in vivo* procedures.

Diagnostic and therapeutic agents that may be attached to a VESPR polypeptide include, but are not limited to, drugs, toxins, radionuclides, chromophores, enzymes that catalyze a colorimetric or fluorometric reaction, and the
10 like, with the particular agent being chosen according to the intended application. Examples of drugs include those used in treating various forms of cancer, e.g., nitrogen mustards such as L-phenylalanine nitrogen mustard or cyclophosphamide, intercalating agents such as cis-diaminodichloroplatinum, antimetabolites such as 5-fluorouracil, vinca alkaloids such as vincristine, and antibiotics such as bleomycin,
15 doxorubicin, daunorubicin, and derivatives thereof. Among the toxins are ricin, abrin, diphtheria toxin, *Pseudomonas aeruginosa* exotoxin A, ribosomal inactivating proteins, mycotoxins such as trichothecenes, and derivatives and fragments (e.g., single chains) thereof. Radionuclides suitable for diagnostic use include, but are not limited to, ^{123}I , ^{131}I , $^{99\text{m}}\text{Tc}$, ^{111}In , and ^{76}Br . Radionuclides suitable for
20 therapeutic use include, but are not limited to, ^{131}I , ^{211}At , ^{77}Br , ^{186}Re , ^{188}Re , ^{212}Pb , ^{212}Bi , ^{109}Pd , ^{64}Cu , and ^{67}Cu .

Such agents may be attached to the semaphorin receptor by any suitable conventional procedure. VESPR, being a protein, comprises functional groups on amino acid side chains that can be reacted with functional groups on a desired agent
25 to form covalent bonds, for example. Alternatively, the protein or agent may be derivatized to generate or attach a desired reactive functional group. The derivatization may involve attachment of one of the bifunctional coupling reagents available for attaching various molecules to proteins (Pierce Chemical Company, Rockford, Illinois). A number of techniques for radiolabeling proteins are known.
30 Radionuclide metals may be attached to the receptor by using a suitable bifunctional chelating agent, for example.

Conjugates comprising VESPR and a suitable diagnostic or therapeutic agent (preferably covalently linked) are thus prepared. The conjugates are administered or otherwise employed in an amount appropriate for the particular application.

35 Another use of the VESPR of the present invention is as a research tool for studying the role that the receptor, in conjunction with semaphorins, may play in immune regulation and viral infection. The VESPR polypeptides of the present invention also may be employed in *in vitro* assays for detection of semaphorin to which it binds or VESPR, or the interactions thereof.

5 As described in Example 16 semaphorins interact with their membrane bound receptors of the present invention to synergize with interferon and *Staphylococcus aureus* (type C) (SAC) in the production of IL-12 from dendritic cells. The use of VESPR and its semaphorin ligand to induce IL-12 production promotes natural killer cell and T cell production and induces cytokine production (primarily γ -interferon).
10 IL-12 and IL-12 induced γ interferon production favors Th1 cell differentiation, and downregulates the production of cytokines associated with Th2 cell differentiation. IL-12 is known to act as both a proinflammatory cytokine and an immunomodulator. Thus, a soluble VESPR can be used to antagonize IL-12 production and downregulate an organism's Th1 cell differentiation. Similarly, a soluble VESPR can be used to
15 promote production of cytokines associated with Th2 cell differentiation, thus discouraging proinflammatory activity. Also, VESPR in combination with its semaphorin ligand can be used to boost IL-12 production in combination with vaccination for those pathogens against which cellular immunity are effective. In this manner the enhanced amount of IL-12 acts as an adjuvant in the vaccination to induce
20 a more persistent Th1-type immunological memory.

 Furthermore, it is known that administration of IL-12 to tumor bearing animals results in tumor regression and the establishment of a tumor-specific immune response. Thus, using a semaphorin ligand to bind with VESPR in order to enhance or promote IL-12 can induce a curative immune response against aggressive
25 micrometastasizing tumors.

 Additionally, as described in example 18, receptors of the present invention bind with their semaphorin ligands to increase CD54 expression on monocytes. This observation suggests that the semaphorin/semaphorin receptor interaction mediate cellular activation that contributes to the proinflammatory activity typically associated
30 with monocyte activation. Such activity includes increased phagocytosis, pinocytosis, nitric oxide production and cytokine production. To antagonize or reverse the proinflammatory activity resulting from the interaction between the semaphorin ligand and its membrane bound receptor, a pharmaceutical composition containing a therapeutically effect amount of a soluble VESPR of the present
35 invention can be administered parenterally to an organism. The soluble VESPR binds with the semaphorin ligand thus preventing the ligand from binding with a membrane bound receptor and contributing to the proinflammatory activity. A therapeutically effect amount of VESPR is an amount sufficient to antagonize proinflammatory activity.

5 Notwithstanding the increased expression of CD54 on monocytes, microphysiometer data indicate that cellular signaling through VESPR does not activate the cell in a classical immunological sense. More particularly, microphysiometer data suggests that when VESPR binds with its ligand a decrease in the rate of change of the pH of the medium results. This is the opposite of that which
10 occurs during cell activation. Such a decrease is experienced with drugs that inhibit protein kinases in the cell, or with viral infections which can paralyze the cell metabolically. Suppressive signals that are delivered through a VESPR can be antagonized by the administration of soluble forms of VESPR. Such soluble forms effectively bind the VESPR binding partner and antagonize the suppressive signaling,
15 thus preventing the inactivation status of the cell. Alternatively, and in accordance with the present invention, anti-VESPR antibodies that signal can be used as a therapeutic to treat autoimmune diseases in which inflammation is a result of presentation of self antigens to T cells. More particularly, such anti-VESPR antibodies can be targeted to be taken up by antigen presenting cells, and, like protein
20 kinase inhibitors, inactivate the antigen presenting cells. The autoimmunity is cured because the antigen presenting cells responsible for the inflammation have become poor antigen presenters.

 Semaphorin ligands binding with VESPR to downregulate expression of MHC Class II molecules and CD86, a co-stimulatory molecule, on dendritic cells,
25 cultured with GM-CSF and IL-4 (see example 17) suggests that the interaction between semaphorin ligands and the receptors of the present invention are associated with the immune suppression of mature dendritic cells. To antagonize or reverse the immunosuppression activity resulting from the interaction between the semaphorin ligand and its membrane bound receptor, a pharmaceutical composition containing a
30 therapeutically effective amount of a soluble VESPR of the present invention can be administered parenterally to an organism. The soluble VESPR binds with the semaphorin ligand thus preventing the ligand from binding with a membrane bound receptor and contributing to the immunosuppression activity. Alternatively, in patients or organisms that suffer from the effects of chronic inflammation,
35 administering appropriate semaphorin ligands will contribute to suppressing the proinflammatory activity of differentiated macrophages.

 Data indicate that a VESPR ligand is found on T cells and VESPR is involved in the migration of dendritic cells from the T cell zones of lymph nodes. Additionally, data suggests that VESPR is involved in shutting down the T cell
40 immune response once a pathogen has been cleared.

5 VESPR polypeptides of the invention can be formulated according to known methods used to prepare pharmaceutically useful compositions. VESPR can be combined in admixture, either as the sole active material or with other known active materials, with pharmaceutically suitable diluents (e.g., Tris-HCl, acetate, phosphate),
10 preservatives (e.g., Thimerosal, benzyl alcohol, parabens), emulsifiers, solubilizers, adjuvants and/or carriers. Suitable carriers and their formulations are described in Remington's Pharmaceutical Sciences, 16th ed. 1980, Mack Publishing Co. In addition, such compositions can contain VESPR polypeptide complexed with polyethylene glycol (PEG), metal ions, or incorporated into polymeric compounds such as polyacetic acid, polyglycolic acid, hydrogels, etc., or incorporated into
15 liposomes, microemulsions, micelles, unilamellar or multilamellar vesicles, erythrocyte ghosts or spheroblasts. Such compositions will influence the physical state, solubility, stability, rate of *in vivo* release, and rate of *in vivo* clearance of VESPR. VESPR polypeptide can also be conjugated to antibodies against tissue-specific receptors, ligands or antigens, or coupled to ligands of tissue-specific
20 receptors.

 VESPR polypeptides can be administered topically, parenterally, or by inhalation. The term "parenteral" includes subcutaneous injections, intravenous, intramuscular, intracisternal injection, or infusion techniques. These compositions will typically contain an effective amount of the VESPR, alone or in combination
25 with an effective amount of any other active material. Such dosages and desired drug concentrations contained in the compositions may vary depending upon many factors, including the intended use, patient's body weight and age, and route of administration. Preliminary doses can be determined according to animal tests, and the scaling of dosages for human administration can be performed according to art-accepted
30 practices.

 VESPR polypeptides may exist as oligomers, such as covalently-linked or non-covalently-linked dimers or trimers. Oligomers may be linked by disulfide bonds formed between cysteine residues on different VESPR molecules. In one embodiment of the invention, a VESPR dimer is created by fusing VESPR to the Fc
35 region of an antibody (e.g., IgG1) in a manner that does not interfere with binding of VESPR to a semaphorin ligand-binding domain. The Fc polypeptide preferably is fused to the C-terminus of a soluble VESPR (comprising only the ligand-binding domain). General preparation of fusion proteins comprising heterologous polypeptides fused to various portions of antibody-derived polypeptides (including
40 the Fc domain) has been described, e.g., by Ashkenazi et al. (*PNAS USA* 88:10535,

5 1991) and Byrn et al. (*Nature* 344:677, 1990), hereby incorporated by reference. A gene fusion encoding the VESPR:Fc fusion protein is inserted into an appropriate expression vector. VESPR:Fc fusion proteins are allowed to assemble much like antibody molecules, whereupon interchain disulfide bonds form between Fc polypeptides, yielding divalent . If fusion proteins are made with both heavy and
10 light chains of an antibody, it is possible to form a VESPR oligomer with as many as four VESPR extracellular regions. Alternatively, one can link two soluble VESPR domains with a peptide linker.

Suitable host cells for expression of VESPR polypeptides include prokaryotes, yeast or higher eukaryotic cells. Appropriate cloning and expression vectors for use
15 with bacterial, fungal, yeast, and mammalian cellular hosts are described, for example, in Pouwels et al. *Cloning Vectors: A Laboratory Manual*, Elsevier, New York, (1985). Cell-free translation systems could also be employed to produce VESPR polypeptides using RNAs derived from DNA constructs disclosed herein.

Prokaryotes include gram negative or gram positive organisms, for example,
20 *E. coli* or *Bacillus*. Suitable prokaryotic host cells for transformation include, for example, *E. coli*, *Bacillus subtilis*, *Salmonella typhimurium*, and various other species within the genera *Pseudomonas*, *Streptomyces*, and *Staphylococcus*. In a prokaryotic host cell, such as *E. coli*, a VESPR polypeptide may include an N-terminal methionine residue to facilitate expression of the recombinant polypeptide in the
25 prokaryotic host cell. The N-terminal methionine may be cleaved from the expressed recombinant VESPR polypeptide.

VESPR polypeptides may be expressed in yeast host cells, preferably from the *Saccharomyces* genus (e.g., *S. cerevisiae*). Other genera of yeast, such as *Pichia*, *K. lactis* or *Kluyveromyces*, may also be employed. Yeast vectors will often contain an
30 origin of replication sequence from a 2 μ yeast plasmid, an autonomously replicating sequence (ARS), a promoter region, sequences for polyadenylation, sequences for transcription termination, and a selectable marker gene. Suitable promoter sequences for yeast vectors include, among others, promoters for metallothionein, 3-phosphoglycerate kinase (Hitzeman et al., *J. Biol. Chem.* 255:2073, 1980) or other glycolytic enzymes (Hess et al., *J. Adv. Enzyme Reg.* 7:149, 1968; and Holland et al.,
35 *Biochem.* 17:4900, 1978), such as enolase, glyceraldehyde-3-phosphate dehydrogenase, hexokinase, pyruvate decarboxylase, phosphofructokinase, glucose-6-phosphate isomerase, 3-phosphoglycerate mutase, pyruvate kinase, triosephosphate isomerase, phosphoglucose isomerase, and glucokinase. Other suitable vectors and
40 promoters for use in yeast expression are further described in Hitzeman, EPA-73,657

5 or in Flier et. al., *Gene*, 107:285-195 (1991); and van den Berg et. al.,
Bio/Technology, 8:135-139 (1990). Another alternative is the glucose-repressible
ADH2 promoter described by Russell et al. (*J. Biol. Chem.* 258:2674, 1982) and
Beier et al. (*Nature* 300:724, 1982). Shuttle vectors replicable in both yeast and *E.*
10 *coli* may be constructed by inserting DNA sequences from pBR322 for selection and
replication in *E. coli* (Amp^r gene and origin of replication) into the above-described
yeast vectors.

The yeast α -factor leader sequence may be employed to direct secretion of the
VESPR polypeptide. The α -factor leader sequence is often inserted between the
promoter sequence and the structural gene sequence. See, e.g., Kurjan et al., *Cell*
15 30:933, 1982; Bitter et al., *Proc. Natl. Acad. Sci. USA* 81:5330, 1984; U. S. Patent
4,546,082; and EP 324,274. Other leader sequences suitable for facilitating secretion
of recombinant polypeptides from yeast hosts are known to those of skill in the art. A
leader sequence may be modified near its 3' end to contain one or more restriction
sites. This will facilitate fusion of the leader sequence to the structural gene.

20 Yeast transformation protocols are known to those of skill in the art. One
such protocol is described by Hinnen et al., *Proc. Natl. Acad. Sci. USA* 75:1929,
1978. The Hinnen et al. protocol selects for Trp⁺ transformants in a selective
medium, wherein the selective medium consists of 0.67% yeast nitrogen base, 0.5%
casamino acids, 2% glucose, 10 μ g/ml adenine and 20 μ g/ml uracil.

25 Yeast host cells transformed by vectors containing ADH2 promoter sequence
may be grown for inducing expression in a "rich" medium. An example of a rich
medium is one consisting of 1% yeast extract, 2% peptone, and 1% glucose
supplemented with 80 μ g/ml adenine and 80 μ g/ml uracil. Depression of the ADH2
promoter occurs when glucose is exhausted from the medium.

30 Mammalian or insect host cell culture systems could also be employed to
express recombinant VESPR polypeptides. Baculovirus systems for production of
heterologous proteins in insect cells are reviewed by Luckow and Summers,
Bio/Technology 6:47 (1988). Established cell lines of mammalian origin also may be
employed. Examples of suitable mammalian host cell lines include the COS-7 line of
35 monkey kidney cells (ATCC CRL 1651) (Gluzman et al., *Cell* 23:175, 1981), L cells,
C127 cells, 3T3 cells (ATCC CCL 163), Chinese hamster ovary (CHO) cells, HeLa
cells, and BHK (ATCC CRL 10) cell lines, and the CV-1/EBNA-1 cell line derived
from the African green monkey kidney cell line CV1 (ATCC CCL 70) as described by
McMahan et al. (*EMBO J.* 10: 2821, 1991).

5 Transcriptional and translational control sequences for mammalian host cell expression vectors may be excised from viral genomes. Commonly used promoter sequences and enhancer sequences are derived from Polyoma virus, Adenovirus 2, Simian Virus 40 (SV40), and human cytomegalovirus. DNA sequences derived from the SV40 viral genome, for example, SV40 origin, early and late promoter, enhancer, 10 splice, and polyadenylation sites may be used to provide other genetic elements for expression of a structural gene sequence in a mammalian host cell. Viral early and late promoters are particularly useful because both are easily obtained from a viral genome as a fragment which may also contain a viral origin of replication (Fiers et al., *Nature* 273:113, 1978). Smaller or larger SV40 fragments may also be used, 15 provided the approximately 250 bp sequence extending from the *Hind* III site toward the *Bgl* I site located in the SV40 viral origin of replication site is included.

Exemplary expression vectors for use in mammalian host cells can be constructed as disclosed by Okayama and Berg (*Mol. Cell. Biol.* 3:280, 1983). A useful system for stable high level expression of mammalian cDNAs in C127 murine mammary epithelial cells can be constructed substantially as described by Cosman et al. (*Mol. Immunol.* 23:935, 1986). A useful high expression vector, PMLSV N1/N4, described by Cosman et al., *Nature* 312:768, 1984 has been deposited as ATCC 39890. Additional useful mammalian expression vectors are described in EP-A-0367566, and in U.S. Patent Application Serial No. 07/701,415, filed May 16, 1991, 25 incorporated by reference herein. The vectors may be derived from retroviruses. In place of the native signal sequence, and in addition to an initiator methionine, a heterologous signal sequence may be added, such as the signal sequence for IL-7 described in United States Patent 4,965,195; the signal sequence for IL-2 receptor described in Cosman et al., *Nature* 312:768 (1984); the IL-4 signal peptide described in EP 367,566; the type I IL-1 receptor signal peptide described in U.S. Patent 30 4,968,607; and the type II IL-1 receptor signal peptide described in EP 460,846.

VESPR polypeptides as isolated, purified or homogeneous proteins according to the invention may be produced by recombinant expression systems as described above or purified from naturally occurring cells. VESPR can be purified to 35 substantial homogeneity, as indicated by a single protein band upon analysis by SDS-polyacrylamide gel electrophoresis (SDS-PAGE).

One process for producing VESPR comprises culturing a host cell transformed with an expression vector comprising a DNA sequence that encodes VESPR polypeptide under conditions sufficient to promote expression of VESPR 40 polypeptide. The receptor is then recovered from culture medium or cell extracts,

5 depending upon the expression system employed. As is known to the skilled artisan, procedures for purifying a recombinant protein will vary according to such factors as the type of host cells employed and whether or not the recombinant protein is secreted into the culture medium.

10 For example, when expression systems that secrete the recombinant protein are employed, the culture medium first may be concentrated using a commercially available protein concentration filter, for example, an Amicon or Millipore Pellicon ultrafiltration unit. Following the concentration step, the concentrate can be applied to a purification matrix such as a gel filtration medium. Alternatively, an anion exchange resin can be employed, for example, a matrix or substrate having pendant
15 diethylaminoethyl (DEAE) groups. The matrices can be acrylamide, agarose, dextran, cellulose or other types commonly employed in protein purification. Alternatively, a cation exchange step can be employed. Suitable cation exchangers include various insoluble matrices comprising sulfopropyl or carboxymethyl groups. Sulfopropyl groups are preferred. Finally, one or more reversed-phase high
20 performance liquid chromatography (RP-HPLC) steps employing hydrophobic RP-HPLC media, (e.g., silica gel having pendant methyl or other aliphatic groups) can be employed to further purify VESPR polypeptide. Some or all of the foregoing purification steps, in various combinations, are well known and can be employed to provide a substantially homogeneous recombinant protein.

25 It is possible to utilize an affinity column comprising the receptor-binding domain of a semaphorin that binds VESPR to affinity-purify expressed VESPR polypeptides. VESPR polypeptides can be removed from an affinity column using conventional techniques, e.g., in a high salt elution buffer and then dialyzed into a lower salt buffer for use or by changing pH or other components depending on the
30 affinity matrix utilized. Alternatively, the affinity column may comprise an antibody that binds VESPR. Example 20 describes a procedure for employing VESPR of the invention to generate monoclonal antibodies directed against VESPR

35 Recombinant protein produced in bacterial culture can be isolated by initial disruption of the host cells, centrifugation, extraction from cell pellets if an insoluble polypeptide, or from the supernatant fluid if a soluble polypeptide, followed by one or more concentration, salting-out, ion exchange, affinity purification or size exclusion chromatography steps. Finally, RP-HPLC can be employed for final purification steps. Microbial cells can be disrupted by any convenient method, including freeze-thaw cycling, sonication, mechanical disruption, or use of cell lysing agents.

5 Transformed yeast host cells are preferably employed to express VESPR as a secreted polypeptide in order to simplify purification. Secreted recombinant polypeptide from a yeast host cell fermentation can be purified by methods analogous to those disclosed by Urdal et al. (*J. Chromatog.* 296:171, 1984). Urdal et al. describe two sequential, reversed-phase HPLC steps for purification of recombinant
10 human IL-2 on a preparative HPLC column.

Useful fragments of the VESPR nucleic acids include antisense or sense oligonucleotides comprising a single-stranded nucleic acid sequence (either RNA or DNA) capable of binding to target VESPR mRNA (sense) or VESPR DNA (antisense) sequences. Antisense or sense oligonucleotides, according to the present
15 invention, comprise a fragment of the coding region of VESPR cDNA. Such a fragment generally comprises at least about 14 nucleotides, preferably from about 14 to about 30 nucleotides. The ability to derive an antisense or a sense oligonucleotide, based upon a cDNA sequence encoding a given protein is described in, for example, Stein and Cohen (*Cancer Res.* 48:2659, 1988) and van der Krol et al. (*BioTechniques*
20 6:958, 1988).

Binding of antisense or sense oligonucleotides to target nucleic acid sequences results in the formation of duplexes that block transcription or translation of the target sequence by one of several means, including enhanced degradation of the duplexes, premature termination of transcription or translation, or by other means. The
25 antisense oligonucleotides thus may be used to block expression of VESPR proteins. Antisense or sense oligonucleotides further comprise oligonucleotides having modified sugar-phosphodiester backbones (or other sugar linkages, such as those described in WO91/06629) and wherein such sugar linkages are resistant to endogenous nucleases. Such oligonucleotides with resistant sugar linkages are stable
30 *in vivo* (i.e., capable of resisting enzymatic degradation) but retain sequence specificity to be able to bind to target nucleotide sequences. Other examples of sense or antisense oligonucleotides include those oligonucleotides which are covalently linked to organic moieties, such as those described in WO 90/10448, and other moieties that increases affinity of the oligonucleotide for a target nucleic acid
35 sequence, such as poly-(L-lysine). Further still, intercalating agents, such as ellipticine, and alkylating agents or metal complexes may be attached to sense or antisense oligonucleotides to modify binding specificities of the antisense or sense oligonucleotide for the target nucleotide sequence.

Antisense or sense oligonucleotides may be introduced into a cell containing
40 the target nucleic acid sequence by any gene transfer method, including, for example,

5 CaPO₄-mediated DNA transfection, electroporation, or by using gene transfer vectors such as Epstein-Barr virus. Antisense or sense oligonucleotides are preferably introduced into a cell containing the target nucleic acid sequence by insertion of the antisense or sense oligonucleotide into a suitable retroviral vector, then contacting the cell with the retrovirus vector containing the inserted sequence, either *in vivo* or *ex vivo*. Suitable retroviral vectors include, but are not limited to, those derived from the murine retrovirus M-MuLV, N2 (a retrovirus derived from M-MuLV), or the double copy vectors designated DCT5A, DCT5B and DCT5C (see PCT Application US 90/02656).

15 Sense or antisense oligonucleotides also may be introduced into a cell containing the target nucleotide sequence by formation of a conjugate with a ligand binding molecule, as described in WO 91/04753. Suitable ligand binding molecules include, but are not limited to, cell surface receptors, growth factors, other cytokines, or other ligands that bind to cell surface receptors. Preferably, conjugation of the ligand binding molecule does not substantially interfere with the ability of the ligand binding molecule to bind to its corresponding molecule or receptor, or block entry of the sense or antisense oligonucleotide or its conjugated version into the cell.

20 Alternatively, a sense or an antisense oligonucleotide may be introduced into a cell containing the target nucleic acid sequence by formation of an oligonucleotide-lipid complex, as described in WO 90/10448. The sense or antisense oligonucleotide-lipid complex is preferably dissociated within the cell by an endogenous lipase.

In addition to the above, the following examples are provided to illustrate particular embodiments and not to limit the scope of the invention.

EXAMPLE 1

Preparing an Ectromelia Semaphorin/Fc Fusion Protein

30 The following describes preparation of an Ectromelia Semaphorin A39R/immunoglobulin fusion protein (A39R/Fc). The process included preparing a DNA construct that encodes the fusion protein, transfecting a cell line with the DNA construct, and harvesting supernatants from the transfected cells. The A39R/Fc fusion protein was used as described in Examples 3, 4, 5 and 6 to study VESPR binding characteristics and isolate VESPR.

35 DNA encoding A39R semaphorin was isolated and amplified from genomic Ectromelia virus DNA using PCR techniques and synthesized oligonucleotide primers whose sequences were based on published A39R sequences in the Copenhagen strain of Vaccinia Virus. The Copenhagen strain A39R DNA sequence is described in

5 Goebel, S.J. et al. *Virology* 179:247, 1990. The isolated Ectromelia A39R DNA is presented in SEQ ID NO:7 and the protein encoded by the DNA is presented in SEQ ID NO:8. The upstream oligonucleotide primer introduced an SpeI site upstream of amino acid 15 of the A39R polypeptide. A downstream oligonucleotide primer introduced a NotI site downstream of the termination codon of Ectromelia A39R,
10 after amino acid 399. The primer sequences were as follows:

Upstream SpeI primer:

TGTCACTAGT ATCGAATGGC ATAAGTTTGA A (SEQ ID NO:3)
Spe I A39R DNA

15

Downstream NotI primer:

GACAGCGGCC GCCTATTACA TTTTAAGTAT TTT (SEQ ID NO:4)
Not I A39R DNA

20 A Bgl II to Nsi I restriction fragment containing a mutein human Fc region of immunoglobulin as described by Baum et al. *Cir. Sh.* 44:30 (1994) was ligated into an expression vector (pDC304) containing a murine IL-7 signal peptide and a FLAG™ octapeptide as described in U.S. Patent No. 5,011,912. The PCR amplified DNA encoding amino acids 15-399 of Ectromelia A39R was then ligated into the
25 expression vector containing the mutein human Fc region, the murine IL-7 signal peptide and FLAG™ peptide, in a two way ligation. The resulting DNA construct was transfected into the monkey kidney cell lines CV-1/EBNA (with co-transfection of psv3neo). After 7 days of culture in medium containing 0.5% low immunoglobulin bovine serum, a solution of 0.2% azide was added to the supernatant
30 and the supernatant was filtered through a 0.22 µm filter. Then approximately 1 L of culture supernatant was passed through a BioCad Protein A HPLC protein purification system using a 4.6 x 100 mm Protein A column (POROS 20A from PerSeptive Biosystems) at 10 mL/min. The Protein A column binds the Fc Portion of the fusion protein in the supernatant, immobilizing the fusion protein and allowing
35 other components of the supernatant to pass through the column. The column was washed with 30 mL of PBS solution and bound fusion protein was eluted from the HPLC column with citric acid adjusted to pH 3.0. Eluted purified fusion protein was neutralized as it eluted using 1M HEPES solution at pH 7.4.

40

5

EXAMPLE 2**Preparing An Ectromelia Semaphorin/polyHis Fusion Protein**

The following describes preparation of an Ectromelia A39R/polyHis fusion protein (A39R/polyHis). The process included preparing a DNA construct that encodes the fusion protein, transfecting a cell line with the DNA construct, and
10 harvesting supernatants from the transfected cells.

DNA encoding Ectromelia A39R (amino acids 1-399 of A39R ORF, SEQ ID NO:8) was isolated and amplified from genomic Ectromelia virus DNA using PCR techniques and synthesized oligonucleotide primers. The primers added a Not 1 site at the 5' end and the motif Gly-Ser-6xHIS at the 3' end for use in purification
15 processes. After the Gly-Ser-6xHIS motif the primers added an in-frame termination codon and a Bgl 2 site. The PCR product was cut and cloned in pDC409 expression vector (McMahon et al., *EMBO J.* 10:2821,1991).

The resulting DNA construct was transiently transfected into the monkey cell line COS-1 (ATCC CRL-1650). Following a 7 day culture in medium containing
20 0.5% low immunoglobulin bovine serum, cell supernatants were harvested and a solution of 0.2% sodium azide was added to the supernatants. The supernatants were filtered through a 0.22 µm filter, concentrated 10 fold with a prep scale concentrator (Millipore; Bedford, MA) and purified on a BioCad HPLC protein purification equipped with a Nickel NTA Superflow self pack resin column (Qiagen, Santa
25 Clarita, CA). After the supernatant passed through the column, the column was washed with Buffer A (20mM NaPO₄, pH7.4; 300mMNaCl; 50mM Imidazole). Bound protein was then eluted from the column using a gradient elution techniques. Fractions containing protein were collected and analyzed on a 4-20% SDS-PAGE reducing gel. Peaks containing A39R/polyHis fusion protein were pooled,
30 concentrated 2 fold, and then dialyzed in PBS. The resulting A39R/polyHis fusion protein was then filtered through a 0.22µm sterile filter.

EXAMPLE 3**Screening Cell Lines for Binding to A39R**

The A39R/Fc fusion protein prepared as described in Example 1 was used to
35 screen cell lines for binding using quantitative binding studies according to standard flow cytometry methodologies. For each cell line screened, the procedure involved incubating approximately 100,000 of the cells blocked with 2% FCS (fetal calf serum), 5% normal goat serum and 5% rabbit serum in PBS for 1 hour. Then the
40 blocked cells were incubated with 5 µg/mL of A39R/Fc fusion protein in 2% FCS,

- 5 5% goat serum and 5% rabbit serum in PBS. Following the incubation the sample was washed 2 times with FACS buffer (2% FCS in PBS) and then treated with mouse anti human Fc/biotin (purchased from Jackson Research) and SAPE (streptavidin-phycoerythrin purchased from Molecular Probes). This treatment causes the antihuman Fc/biotin to bind to any bound A39R/Fc and the SAPE to bind to the anti-
- 10 human Fc/biotin resulting in a fluorescent identifying label on A39R/Fc which is bound to cells. The cells were analyzed for any bound protein using fluorescent detection flow cytometry. The results indicated the A39R semaphorin binds well to human NK cells, murine splenic B cells, human PB T cells, human T, B, erythroid, lymphoid and myeloid precursor cells, fibroblasts and epithelial lineage. Table I
- 15 details the results of the flow cytometry studies. A "+" indicates that binding was detected between the cell surface and A39R. A "-" indicates that no binding was detected between the cell surface and A39R.

TABLE I

20	<u>Cell Line</u>	<u>A39R Binding Result</u>
	Namalwa (B cell-like lymphoma - human)	+
	CB23 (Human Cord Blood B Cell Line)	+
	EU-1 (Human pre B Cell Line)	+
	MP-1 (Human B Cell Lymphoma)	+
25	PB B (Human Peripheral Blood B Cells)	+
	Mouse Splenic B Cells	+
	Mouse Splenic B Cells + CD40L	+
	U937 (Human Monocyte-Type Cell)	+
	HSB2 (Human T Cell Line)	+
30	K299 (Non Hodgkin's Lymphoma)	+
	TE71 (Mouse Thymic Epithelium)	+
	IEC18 (Rat Intestinal Epithelium)	+
	IMTLH (Human Bone Marrow Derived Stroma)	+
	W126 (Human Lung Epithelium)	+
35	PL-1 (Human T-Cell Clone)	+
	VK-1 Human T-Cell Clone	+
	Primary Peripheral Blood T Cells	+
	Primary Human NK Cells	+
	RAJI (Burkitt's Lymphoma)	-
40	KG1 (Human myeloid Cell Line)	-

5	THP-1	(Human Promonocytic Cell Line)	+
	MC6	(Mouse Mast Cell)	-
	EL4	(Mouse Thymoma)	-
	BeWo	(Chorio Carcinoma)	-
	Primary Mouse Dendritic Cells		+
10	Primary Human Dendritic Cells		+

EXAMPLE 4

Identifying A Putative Semaphorin Receptor

CB23 cells (human cord blood B cell line) and human PB T cells that tested positive for binding to A39R were tested for expression of a putative receptor and to determine if any receptor is expressed as a membrane bound molecule, soluble molecule, or both. Broadly, the analyses involved radiolabeling CB23 and human PB T cell surfaces, harvesting and treating cell supernatants and lysates with an A39R/Fc fusion protein to precipitate any putative receptor, and then visualizing an immunoprecipitate on an electrophoretic gel.

In particular, the procedure involved first radiolabeling approximately 1×10^7 CB23 or PB T cells with [125 I] as described by Benjamin et al.; *Blood* 75,:2017-2023 (1990). Cultured cell supernatants were harvested and clarified by centrifugation at 14,000 rpm for 30 minutes. Cell lysates were generated by incubating the cells on ice for 30 minutes in 1 mm L phosphate-buffered saline with 1% Triton-X 100 containing protease inhibitors phenylmethylsulfonyl fluoride, Pepstatin-A, and Leupeptin. The lysates were clarified by centrifugation at 14,000 rpm for 30 minutes. In order to precipitate any receptor present in the lysate and/or supernatant, 200 μ L of the cell supernatant or lysate was incubated with 2 μ g of A39R/Fc fusion protein prepared as described in Example 1. The incubation was carried out for 1 hour with gentle rocking at 4°C. An Fc protein control sample was prepared and incubation in the same manner. Following the incubation, Protein-A Sepharose beads (#17-0780-01 Pharmacia Biotech Inc., Piscataway, NJ) were added to the lysates and supernatants and the mixture was incubated for 1 hour with gentle rocking at 4°C. The beads were washed extensively with a PBS 1% Triton-X 100 solution. Bound protein was eluted and analyzed by SDS PAGE. Protein bands were visualized by autoradiography and a single, approximately 200K Da band was found to bind to A39R/Fc but not to the control Fc Protein. The semaphorin receptor was present in cell lysate and cell supernatant, confirming its expression as membrane bound protein and as a secreted soluble protein.

5

Example 5

Isolating and Sequencing a Semaphorin Receptor

The A39R/Fc fusion protein, prepared as described in Example 1, was used to isolate a human semaphorin receptor polypeptide and a procedure for the isolated polypeptide purification was confirmed. The semaphorin receptor was isolated by suspending CB23 cell pellets in a solution of protease inhibitors that included 1 mM each of PMSF, Leupeptin, Aprotinin, Pepstatin A, 10 µg/mL APMSF, and 1 mM EDTA in homogenization buffer (10 mM phosphate, 30 mM NaCl, pH 7.4). The cells were dounce homogenized and layered over a solution of 41% sucrose in homogenization buffer and the spun down in a Beckman SW-28 rotor at 25,000 rpm, at 4°C for 45 minutes. The interphases were collected and diluted in cold homogenization buffer, dounced, and spun. The resulting clean membrane pellets were stored at -80 °C.

Membrane pellets prepared from 240 mLs of packed cells were combined with 100 mLs of an aqueous solution of 20 mM Tris, 150 mM NaCl, the protease inhibitors identified above, 1% Triton X-100 and 0.1 mM of CaCl₂, MgCl₂, and MnCl₂ salts (Buffer A). The suspended pellets were dounced and spun in a SW-28 rotor for 30 minutes at 25,000 rpm at 4°C. The supernatant was placed onto a 100mL wheat germ agglutinin column and allowed to elute at a rate of 1 mL/minute with 10 column volumes of Buffer A. Proteins that were specifically bound to the column were then eluted with Buffer A containing 0.2 M N-acetyl glucosamine.

Fractions testing positive for protein were pooled and incubated with 100 µg of A39R/Fc fusion protein for 1 hour at 4°C. The incubated mixture was run through a sepharose column to remove material that did not specifically bind and then allowed to pass through a 0.5 mL column of Protein A/Sepharose solid support. The Protein A/Sepharose solid support was washed with 20 column volumes of PBS containing 1% Triton X-100 followed by a wash with PBS to wash off any unbound material. Then proteins that were retained on the Protein A/Sepharose column were eluted in a stepwise manner with 0.35 mL fractions of 50 mM citrate at pH 3.0. Fractions that tested positive for protein were combined and concentrated to 50 µL using a 10 kD MWCO Centricon concentrator. Protein in the resulting concentrated sample were reduced and then alkylated using standard DTT and iodoacetic acid procedures. The alkylation proteins were then electrophoresed on an 8% gel. Proteins on the gel were visualized with coomassie-G in 50% MeOH containing 5% acetic acid and then destained in 50% MeOH.

5 The approximately 200 kD band, located by comparison to protein standards, was excised with a razor blade and washed overnight in 100 mM ammonium carbonate. The gel slice was speed evaporated until dry and a 1:10 solution of trypsin in 100 mM ammonium carbonate was added to the dried slide. The slide was incubated at 37°C for 16 hours and then protein in the slice was extracted with 50%
10 acetonitrile with 5% formic acid three times while incubating 30 minutes with each extraction.

 The trypsin digested peptide fragments were lyophilized, reconstituted in 50µL of 0.1% trifluoroacetic acid, and separated by RP-HPLC on a 500 µ id x 25 cm capillary column packed with C-18 reverse phase packing. The HPLC liquid phase
15 was an acetonitrile/water gradient of 10% after 5 minutes, 85% after 105 minutes. Eluting protein was detected at 215 nm. Each protein was collected as it eluted in separate fractions and N-terminal sequence analysis of the peptides in the fraction was performed on a 494 Procise sequencer according to the manufacturer's instruction.

 RP-HPLC fractions, obtained as described above, were dried on a vacuum
20 centrifuge and peptides in the fraction were dissolved in 6µL of 50% methanol containing 0.5% acetic acid. Two microliters (2 µL) of each of the peptide solutions were loaded into nanospray tips (Protein Analysis Company, Odense, Denmark). Data were obtained with a Finnigan TSQ700 triple quadrupole mass spectrometer (San Jose, CA) equipped with a nanospray source. Mass spectra were acquired at unit
25 resolution. For tandem mass spectrometry, the first quadrupole was operated at a resolution sufficient to pass a 3-4 Da wide window, and the third quadrupole was operated at unit resolution. Collision gas was supplied at a pressure of 4 mTorr. Methyl esterification was performed using standard esterification procedures.

 The tandem mass spectrometry analysis of the trypsin generated peptides
30 provided amino acid sequence information for isolated portions of the purified protein. The tandem mass spectral data were used in computer assisted screening of non-redundant protein databases and EST databases using the local SEQUEST algorithm search tool (Eng, J.K et al. J am Soc. Mass 1994). The peptide query sequences GluGluThrProValPheTyrLys corresponding to amino acids 421-428 of
35 SEQ ID NO:2, and AsnIleTyrIleTyrLeuThrAlaGlyLys, corresponding to amino acids 436-445 identified EST No. 248534 (Accession N78220) as containing peptide sequences having 100% identity to the query peptide sequences. The peptide query sequence ThrValLeuPheLeuGlyThrGlyAspGlyGlnLeuLeuLys corresponding to
40 amino acids 388-401 identified EST No. R08946 as containing a 100% identity to the query.

5 The 100% identity between portions of EST 248534 and three peptide fragments of the purified protein strongly suggested that the cDNA contained within EST 248534 represented a portion of the nucleotide sequence for the coding region of the purified protein. A source of semaphorin receptor cDNA was identified using phage library screening methods and PCR primers based upon EST 248534.

10 The oligonucleotide primers had the following nucleotide sequences:

ATCGCATCAT CTACCTTCAT CCATTCCGAC CTG (SEQ ID NO:9)

TAAACACTCC GAACAGGATT TATGTTTATT GCA (SEQ ID NO:10)

15 PCR isolation and amplification methodologies were carried out using a panel of human tissue cDNA phage libraries as templates for the PCR reactions. The PCR reaction mixture included 1 μ L of phage library stock, PCR oligonucleotide primers at 0.3 μ M final concentration, 1x PC2 buffer (Ab Peptides, Inc., St. Louis, MO), 0.2 mM each of dATP, dCTP, dGTP, dTTP (Pharmacia Biotech) 0.2 μ L of a 16:1 mix Klen-Taq/Vent polymerase (Klen-Taq polymerase, Ab Peptides, Inc. and Vent polymerase, New England Biolabs, Beverly, MA) in a 30 μ L final reaction volume. The PCR reaction cycles included one cycle at 98°C for 5 minutes; thirty cycles at 98°C for 45 seconds, thirty cycles at 68°C for 45 seconds; thirty cycles at 72°C for 45 seconds, and 1 cycle at 72°C for 5 minutes using a Robocycler 96 from Stratagene, La Jolla, CA. cDNAs in several libraries were positively identified as containing DNA encoding the purified VESPR protein based upon the appearance of an appropriate sized DNA band in electrophoresed PCR product.

25 Two of the phage libraries, human foreskin fibroblast and human dermal fibroblasts, were chosen for additional analysis. Libraries were plated according to established procedures and probed with a radiolabeled random primer probe derived from a PCR amplification product using EST 248534 as a template. The PCR conditions used to obtain the amplification product were as described above and the probe was generated using Prime-IT II Random Primer Labeling Kit from Stratagene, La Jolla, CA. Approximately 1×10^6 cpm/mL of purified probe was used to probe human foreskin phage libraries on nylon membrane filters overnight at 63°C in a hybridization buffer of 10x Denhardt's solution, 50 mM Tris at pH 7.5, 0.9 M NaCl, 0.01% Sodium Pyrophosphate, 1% sodium dodecyl sulfate, and 200 μ g/mL denatured, fragmented salmon sperm DNA. After probing, the probed membranes were washed once in 6x SSC, 0.1% SDS for 20 minutes, once in 2 x SSC, 0.1% SDS for 20

5 minutes, once in 1x SSC, 0.1% SDS for 20 minutes, and once in 0.1 x SSC, 0.1% SDS for 20 minutes at 63°C. The probed and washed filters were exposed to X-omat AR X-ray film (Eastman-Kodak Corp.) overnight. Four overlapping cDNAs were identified. The overlapping cDNAs, together with the sequenced trypsin digest generated protein fragments were used to complete and confirm the coding sequence of VESPR as shown in SEQ ID NO:1 and the amino acid sequence presented in SEQ ID NO:2.

Example 6
Monoclonal Antibodies to A39R Semaphorin

15 This example illustrates a method for preparing antibodies to A39R semaphorin. Purified A39R/Fc was prepared as described in Example 1 above. The purified protein was used to generate antibodies against A39R semaphorin as described in U.S. Patent 4,411,993. Briefly, mice were immunized at 0, 2 and 6 weeks with 10 µg with A39R/Fc. The primary immunization was prepared with
20 TITERMAX adjuvant, from Vaxcell, Inc., and subsequent immunizations were prepared with incomplete Freund's adjuvant (IFA). At 11 weeks, the mice were IV boosted with 3-4 µg A39R/Fc in PBS. Three days after the IV boost, splenocytes were harvested and fused with an Ag8.653 myeloma fusion partner using 50% aqueous PEG 1500 solution. Hybridoma supernatants were screened for A39R
25 antibodies by dot blot assay against A39R/FC and an irrelevant Fc protein.

Example 7
Northern Blot Analyses for Tissue Expressing Semaphorin Receptor

30 The following describes Northern Blot experiments carried out to identify tissue and cell types that express VESPR polypeptides of the present invention. The results confirm the cell binding results obtained using flow cytometry analysis and the A39R/Fc fusion protein.

As described in Example 5, EST data base searches resulted in the discovery of an EST that was believed to be a partial clone of the VESPR of SEQ ID NO:2 (EST 248534). A riboprobe template was generated using PCR techniques and oligonucleotide primers that were based on nucleotides 1-372 of EST 248534. The upstream and down stream primers that encompasses nucleotides 1-372 of EST 248534 had the following sequences:

40 GCGGGACTCA GAGTCACC (SEQ ID NO:5)

5

GGATCCTAAT ACGACTCACT ATAGGGAGGA AACCACTCCG AAC
(SEQ ID NO:6)

The underlined portion is a T7 site.

The two primers were used to isolate and amplify a PCR product from
10 EST248534 for use in generating a riboprobe. The riboprobe was generated using
Ambion's MAXIscript SP6/T7 kit by combining 3 μ L of RNase free water, 2 μ L 10x
transcription buffer, 1 μ L each of 10mM dATP/dCTP/dGTP, 5 μ L 5'/3' EST 248534
PCR Product, 5 μ L Amersham [α^{32} P]UTP 10mCi/mL, 2 μ L T7 RNA polymerase at
room temperature. The combination was microfuged, spun briefly, and incubated at
15 37°C for 30 minutes. Then 1 μ L DNase was added to the mixture and allowed to
react for 15 minutes at 37°C. The reaction product was passed through two column
volumes of G-25 packing (Boehringer). One microliter (1 μ L) of the riboprobe was
counted in a scintillation counter for 1 minute to determine cpm/mL

Northern blots were generated by fractionating polyadenylated RNA from a
20 variety of cell lines on a 1.2% agarose formaldehyde gel and blotting the RNA onto
Hybond Nylon membranes (Amersham, Arlington Heights, IL). Standard northern
blot generating procedures as described in Maniatis, (*Molecular Cloning: a
Laboratory Manual*, Cold Spring Harbor Lab. Press, 1989) were used. Total RNA
multiple tissue northern blots were purchased commercially (BioChain Institute, Inc.,
25 San Leandro, CA Cat #s 021001, 021002, 021003).

The Northern blots were prehybridized in a 50% formamide hybridization
solution (30 mL 20x SSC, 2 mL 100x Denhardt's reagent, 1 mL of 10 mg/mL
denatured fragmented salmon sperm DNA, 50 mL 100% formamide and 20 mL 10%
SDS. The total RNA blots were pre-hybridized at 42°C for 4 hours and the polyA+
30 RNA blots were pre-hybridized at 63°C for 4 hours. The riboprobe was added to
clean hybridization solution (same as prehybridization solution) at a count of 10⁶
cpm/mL. The prehybridization solution was removed from the blots and the
hybridization solution and riboprobe were added to the blots. The hybridization was
allowed to proceed overnight with gentle shaking. The total RNA blots hybridized at
35 63°C and the polyA+ RNA blots hybridized at 63°C.

The probed total RNA blots were washed once for 30 minutes in 2xSSC
containing 0.05% SDS at 42°C and once for 30 minutes in 2xSSC containing 0.05%
SDS at 55°C; twice for 30 minutes in 0.1xSSC containing 0.1% SDS at 63°C; three
times for 30 minutes in 0.1xSSC containing 0.1%SDS and then exposed to X-ray
40 film. The poly A+ blots were washed once for 30 minutes in a 2xSSC solution

- 5 containing 0.05% SDS at 63°C and once for 30 minutes in 1xSSC containing 0.1% SDS and then exposed to x-ray film.

The results of probing the Northern blots and visualizing the resulting x-ray film for positively binding probes confirm that VESPR is expressed in the same cells as those that showed positive binding in flow cytometry experiments. Hybridizing
10 RNA was detected in MP-1, HFF and CB23 cells. Primary tissues showing positive RNA included heart, brain, lung, spleen and placenta. No RNA was detecting in RAJ1 cells.

Example 8

15 Generating AHV Semaphorin Fc Fusion Protein

The following describes preparing an AHV Semaphorin/immunoglobulin fusion protein (AHVSema/Fc). The process included preparing a DNA construct that encodes the fusion protein, transfecting a cell line with the DNA construct, and harvesting supernatants from the transfected cells.

- 20 DNA encoding AHV-Sema is described in Ensser et al. *J.Gen. Vir.* 76:1063-1067, 1995. DNA encoding AHV-Sema amino acids 70-653 was isolated and amplified from Alcelaphine herpesvirus DNA strain WC11 (Plowright, W. et al. *Nature* 188:1167-1169, 1960) using PCR techniques and synthesized oligonucleotide primers whose sequences were based on the published AHV-Sema sequence. The
25 upstream oligonucleotide primer introduced a Spe 1 site. A downstream oligonucleotide primer introduced a Not 1 site downstream of the termination codon. The general method used to isolate the soluble AHVSema is described in Spriggs et al., *J. Virology*, 70:5557 (1996).

- A restriction fragment containing a mutein human Fc region of
30 immunoglobulin as described by Goodwin et al. *Cell* 73, 447-456, 1993 was ligated into an expression vector (pDC409) containing a murine IL-7 signal peptide and a FLAG™ octapeptide as described in U.S. Patent No. 5,011,912. The PCR amplified AHVSema DNA encoding was then ligated into the expression vector containing the mutein human Fc region, the murine IL-7 signal peptide and FLAG™ peptide, in a
35 two way ligation. The resulting DNA construct was transfected into the monkey kidney cell lines CV-1/EBNA (with co-transfection of pSV3neo). After 7 days of culture in medium containing 0.5% low immunoglobulin bovine serum, a solution of 0.2% azide was added to the supernatant and the supernatant was filtered through a 0.22 µm filter. Then approximately 1 L of culture supernatant was passed through a
40 BioCad Protein A HPLC protein purification system using a 4.6 x 100 mm Protein A

5 column (POROS 20A from PerSeptive Biosystems) at 10 mL/min. The Protein A column binds the Fc Portion of the fusion protein in the supernatant, immobilizing the fusion protein and allowing other components of the supernatant to pass through the column. The column was washed with 30 mL of PBS solution and bound fusion protein was eluted from the HPLC column with citric acid adjusted to pH 3.0. Eluted
10 purified fusions protein was neutralized as it eluted using 1M HEPES solution at pH 7.4.

Example 9

Expressing Recombinant Semaphorin Receptor

15 Using the semaphorin receptor (VESPR) amino acid sequence of the protein purified as described in Example 5, and information derived from EST database searches and cDNAs obtained using hybridization methodologies with radiolabeled probes, also as described in Example 5, cDNA is generated and cells are transfected with the cDNA to allow expression of recombinant VESPR polypeptide.

20 The cDNA in DC409 expression vector, derived from pDC406, is transfected in CV1/EBNA cells using standard techniques (McMahan et al., *EMBO J.* 10:2821,1991) More particularly, CV1 EBNA cells are plated at a density of 2×10^6 cells per 10 cm dish in 10 mL of Dulbeccos Minimum Essential Medium (medium) supplemented with 10% fetal calf serum. The cells are allowed to adhere overnight at
25 37°C. The medium is replaced with 1.5 mL of medium containing 66.7 μ M chloroquine and a DNA mixture containing 5 μ g of cDNA encoding VESPR. Medium containing 175 μ L and 25 μ L of DEAE dextran is added to the cells. The cells and cDNA are incubated at 37°C for 5 hours. The cDNA mixture is removed and cells are shocked with 1 mL of fresh medium containing 10% DMSO for 2.5 min.
30 The medium is replaced with fresh medium and the cells are grown for at least 3 days.

To recover soluble forms of VESPR, supernatants containing the soluble form are collected and the VESPR protein recovered using HPLC techniques or affinity chromatography techniques. To recover forms of VESPR that are membrane bound,
35 the transfected cells are harvested, fixed in 1% paraformaldehyde, washed and used in their intact form.

Example 10

VESPR Binding Studies

In order to examine the binding characteristics of a receptor polypeptide of the
40 present invention, binding studies were performed by subjecting cells expressing

5 membrane bound VESPR extracellular domain to the slide binding assay described in Goodwin et al. *Cell* 73:447-456, (1993) and Spriggs et al., *J Virol* 70:5557 (1996).

The pDC409 expression vector, derived from pDC406 (McMahon et al., *EMBO J.* 10:2821, 1991) but having a single Bgl 2 was selected for the cloning process. VESPR cDNA, encoding amino acids 19-1100, was subcloned into a
10 pDC409 expression vector through the Sal 1 (5') and Not 1 (3') sites, to form a DNA construct.

CV-1/EBNA cells were transfected via DEAE/Dextran with 2 μ g of a VESPR cDNA (encoding amino acids 19-1100) in pDC409 (Giri et al., *EMBO J* 13:2822, 1994). The transfected cells were cultured for 3 days and the CV-1/EBNA cell
15 monolayers were incubated with 1 μ g/mL of A39R/Fc, AHVsema/Fc, or control Fc protein. Then the incubated cells were washed and incubated with ¹²⁵I-labeled mouse anti-human IgG (Jackson ImmunoResearch, West Grove, PA). After extensive washing, the cells were fixed, dipped in photographic emulsion as described by Gearing et al., *EMBO J* 8:3667-3676 (1989) and developed. Positive binding was
20 determined by the presence of exposed or darkened silver grains overlaying cells expressing VESPR that had bound Fc protein.

EXAMPLE 11

Flow Cytometry and Inhibition Binding Studies

25 The following describes flow cytometric analyses of CB23 cells for binding to A39R/Fc fusion protein (Example 1) and the AHVsema/Fc fusion protein (Example 9). Also described below is a study directed to determining inhibition of the AHVsema and A39R binding with and an excess of A39R/polyHis fusion protein prepared as described in Example 2.

30 The flow cytometric analysis was performed by first incubating about 1x10⁶ CB23 cells on ice for 30 minutes in FACS buffer and containing 3% normal goat serum and 3% normal rabbit serum to block non-specific binding. Portions of A39R/Fc, AHVsema/Fc and a control Fc protein were added at varying concentrations and the incubation was continued for 30 minutes. The cells were
35 washed and then incubated with phycoerythrin-conjugated Fc specific anti-human IgG in FACS buffer. The cells were washed and analyzed on a FACScan from Becton Dickinson, Bedford, MA. The results showed positive binding of AHV semaphorin and the A39R semaphorin.

40 Binding inhibition studies were performed by incubating about 1x10⁶ CB23 cells for 30 minutes on ice in FACS buffer. The A39R/polyHis and control HIS

5 protein were added to different samples at varying concentrations and the incubation continued for another 30 minutes. Then A39R/Fc or AHVsema/Fc were added to the incubated cells at varying concentrations and the incubation was continued for another 30 minutes. The cells were washed and then incubated with phycoerythrin-conjugated Fc specific anti-human IgG in FACS buffers. The cells were washed
10 again and then analyzed on a FACScan. The results demonstrated complete inhibition of A39R and AHVsema using A39R/polyHIS, but not the heterologous HIS containing protein.

Example 12

Human B Cell Aggregation with A39R Semaphorin

15 In order to examine human B cell response to A39R semaphorin, human tonsillar B cells were purified as described in Spriggs et al., *J Exp Med* 176:1543, (1992). An A39R/polyHis fusion protein was prepared as described in Example 2. A solution of A39R/polyHis fusion protein was prepared to a final A39R concentration
20 of 1 µg/mL and the A39R/polyHis fusion protein solution was incubated in *in vitro* cultures of about 10⁵ of the purified B cells. Continuing the incubation for about 24 hours resulted in cellular aggregation. When a 10 fold molar excess of the monoclonal antibody against A39R, prepared as described in Example 6, was added to the fusion protein preparation prior to adding the fusion protein to the cultures, the
25 cell aggregation was blocked. Additionally, when the A39R semaphorin was heat inactivated prior to adding it to the culture, the aggregation was blocked.

This work confirms that VESPR is expressed on B cells and that the interaction between A39R and VESPR results in B cell aggregation. B cell aggregation is indicative of their activation. Activated B cells are known to secrete
30 cytokines, produce antibodies, or become antigen presenting cells.

Example 13

Mouse Dendritic Cells and Macrophage Aggregation with A39R

Semaphorin

35 In order to examine dendritic cell and macrophage response to A39R, mouse cell cultures were brought into contact with A39R semaphorin and the effects of the combination noted. Mouse dendritic cell cultures containing macrophages were obtained by immunizing mice with Flt3-L and cells were isolated and purified as described in Maraskovsky et al., *J Exp Med* 184:1953, (1996).

5 Briefly, female C57Bl/6 mice were injected once daily with a solution of 10 μ g of Flt3L and 1 μ g mouse serum albumin in 100 μ L of PBS for 9-10 consecutive days. After the immunization, single cell suspensions of spleens were prepared by disrupting spleen tissue between frosted glass slides in the presence of NH_4Cl to deplete red blood cells. The remaining cells were incubated with mAb to Thy-1,
10 B220, NK1.1, and TER119, and then incubated with 10% rabbit complement. Then the incubated cells were washed and residual mAb-coated cells were removed using anti-immunoglobulin (Ig)-coated magnetic beads. The remaining enriched cells were cultured or sorted for the various cell populations.

Cells selected for sorting were stained with anti-CD11c and anti-CD11b and
15 sorted for the C and D/E populations as described in Maraskovsky et al., *J Exp Med* 184:1953-1962, 1996.

An A39R/polyHis fusion protein was prepared as described in Example 2. An A39R/polyHis fusion protein solution was incubated in *in vitro* cultures at a final concentration of 1 μ g/mL with about 10^5 of the sorted or depleted mouse cells. Within
20 4-6 hours the cells began to aggregate. When a 10 fold molar excess of the monoclonal antibody against A39R, prepared as described in Example 6, was added to the A39R/polyHis fusion protein preparation prior to adding the fusion protein to the mouse cell cultures, the aggregation was blocked.

This work confirms that VESPR is expressed on dendritic cells and
25 macrophages, and that the interaction between A39R and VESPR results in dendritic cell and macrophage aggregation.

Example 14

A39R Semaphorin Upregulates CD69 Activation Antigen

30 In order to investigate the effects of A39R semaphorin on cultured dendritic cells, mice were injected each day for 9 days with a Flt3-L preparation. Mouse dendritic cells were harvested and then cultured in medium containing 10% FBS and 20 ng/mL GM-CSF for 5 days.

On day 5, 1 μ g/mL of A39R/polyHis fusion protein was added to the culture.
35 On day 6, the cells were stained with diagnostic antibodies. The results of the diagnostic antibody staining experiments showed that CD11c^+ , CD11b^+ cells (dendritic cells) expressed an increased amount of the CD69 activation antigen, thus demonstrating that the interaction of A39R semaphorin and its receptor upregulate CD69 expression.

5 When the fusion protein is inactivated with heat, the fusion protein had no effect on the CD69 antigen. Representative changes in mean fluorescence intensity between unstained and stained cells were from approximately 500 channels to 2500 channels. Again, these results demonstrate significant effects of the interactions between A39R semaphorin and its membrane bound receptor on the regulation of the
10 CD69 activation antigen, a transient and early expressed marker for cell activation.

Example 15

Evaluating the Effect of A39R in the Production of IL-12

 In order to study the role of A39R in the production of IL-12 from mouse
15 spleen cells, mice were immunized with flt3-L and dendritic cells were generated, harvested and purified as described in Example 13.

 Approximately 5×10^5 cells/0.5 mL of purified, unsorted dendritic cells were incubated in modified DMEM media (500 μ L at 1×10^6 /mL) in the presence of one more of the following: 20ng/mL muGM-CSF (Immunex, Seattle, WA), 20ng/mL γ -
20 IFN (Genzyme, Boston, MA), 10 μ g/mL SAC (CalBiochem, La Jolla, CA). Each cell preparation was treated additionally with 1 μ g/mL of A39R/polyHis fusion protein alone or in combination with 1 μ g/mL or 0.1 μ g/mL of muCD40L trimer (Immunex, Seattle, WA). Cultures were incubated in humidified 37C, 10% CO₂-in-air for 16-18 h. After incubation, the viability of each group of cultured cells was determined and
25 supernatants were collected and assayed for muIL12 (P70) using an ELISA assay kit (Genzyme, Boston, MA). MuIL12 levels were calculated by reference to a standard curve constructed with recombinant cytokine.

 ELISA testing demonstrated in particular that A39R interacts with its receptor to synergize with interferon and SAC in the production of IL-12 from unsorted mouse
30 dendritic cells. This *in vivo* IL-12 induction promotes natural killer cell activation and gamma interferon production and contributes to upregulating gamma interferon sensitive cytokines.

Example 16

Testing Effects of A39R on Regulation of MHC Class II and CD86 on Monocytes

 The following experiment describes upregulation of MHC Class II and CD86 by the interaction of A39R with its membrane bound receptor. Peripheral blood from healthy donors was diluted 1:1 in low endotoxin PBS at pH 7.4 and room
40 temperature. Then 35 mLs of the diluted blood was layered over 15 mLs of Isolymp

5 (Gallard and Schlesinger Industries, Inc; Carle Place, NY) and centrifuged at 2200 rpm for 25 minutes at room temperature. The plasma layers was reserved. The PBMC layer was harvested and washed three times to remove the Isolymp. The washed PBMC's were resuspended in X-Vivo 15 serum free media (BioWhittaker, Walkersville, MD) and added to T175 flasks. The flasks had been previously coated with 2% Gelatin (Sigma, St. Louis, MO) and pre-treated for 30 minutes with the reserved plasma layer. The PBMC's were allowed to adhere for 90 minutes at 37°C, 5% CO₂ and then rinsed three times gently with 10 mL washes of low endotoxin PBS. Adhered monocytes were harvested by incubating the cells in Enzyme Free Dissociation Buffer (Gibco, BRL) and washing the cells multiple times in PBS. 15 Monocytes were centrifuged at 2500 rpm for 5 minutes, counted, and set up in 24 well dishes at 5×10^5 cells/well in 1 mL. The cultures were 95% pure.

Purified monocytes were cultured for 7-9 days in the presence of 20 ng/mL GM-CSF and 100 ng/mL IL-4 in order to allow cells to differentiate to a more dendritic cell-like phenotype. On day 7-9, cultures were treated with 1 µg/mL A39R/polyHis or a control polyHis containing protein, and the next day cells and supernatants were harvested for analyses. 20

In flow cytometric experiments for examining monocyte-derived dendritic cell surface markers, cells were stained with conjugated mAbs directed against specific proteins. The staining showed that for a majority of the peripheral blood donors tested, A39R treatment downregulated CD86 and MHC class II expression on these cells. Since CD86 and MHC class II molecules are markers of an enhanced antigen presentation by dendritic cells, their downregulation suggests an immunosuppressive effect of the interaction of A39R with its receptor on this cell population. 25

30

Example 17

Upregulation of CD54

The following describes the effect of the interaction between A39R semaphorin and its receptor on purified monocytes and more particularly, the impact of CD54 expression on monocytes after incubation with a semaphorin. Freshly isolated monocytes were purified from peripheral blood donors as described in Example 16, except that they were held in culture overnight in the presence of A39R/polyHis or control proteins. 35

Following the overnight culture, flow cytometry was performed using the cultured cells and mAbs directed against monocyte specific cell surface markers. In all donors tested, the level of CD54 surface expression was enhanced in the presence 40

5 of A39R, but not in the presence of heat inactivated A39R. Similarly, in cultures containing control proteins CD54 surface expression was not enhanced.

CD54, also known as ICAM-1, is an adhesion molecule whose increased expression is considered to be indicative of cellular activation. These data indicate that promoting the interaction of A39R with its receptor can activate freshly isolated
10 human monocytes.

Example 18

Cytokine Induction from Freshly Isolated Human Monocytes

Freshly isolated human monocytes were purified as described in Example 16,
15 and cultured as described in Example 17. After the overnight incubation with A39R/polyHis, monocyte supernatants were examined for the presence of proinflammatory cytokines. In all donors tested, IL-6 and IL-8 was induced by A39R protein. Heat inactivated A39R and control proteins did not induced IL-6 or IL-8. Additionally, cytokine production was blocked by the inclusion of a mAb directed
20 against A39R.

The results of this experiment demonstrate that A39R, or homologues of this protein, interacting with its receptor, can induce cytokine production by freshly isolated monocytes. Advantageously, soluble forms of VESPR can be used in inhibit the proinflammatory activity of monocytes in response to A39R or its homologues.
25

Example 19

Monocyte Aggregation Studies

In order to examine human monocyte response to the interaction of a semaphorin to its receptor on monocytes, monocytes were purified as described in
30 Example 17 and an A39R/polyHIS fusion protein was prepared as described in Example 2. The fusion protein and purified, cultured monocytes were incubated. Continuing the incubation for 20 hours resulted in monocyte aggregation. In view of the results demonstrated in Example 17, it is suggested that the observed monocyte aggregation occurs as a result of CD54 upregulation. However, other factors may
35 contribute to the aggregation as well.

This work confirms that the semaphorin receptor of the present invention is expressed on monocytes and that the interaction between A39R and VESPR results in monocyte aggregation. Similar to B cells, monocytes aggregation is indicative of their activation.
40

5

Example 20**Monoclonal Antibodies to VESPR**

This example illustrates a method for preparing antibodies to VESPR polypeptides. Purified VESPR polypeptide is prepared as described in Example 10. The purified protein is used to generate antibodies against VESPR as described in U.S. Patent 4,411,993. Briefly, mice are immunized at 0, 2 and 6 weeks with 10 μ g with VESPR. The primary immunization is prepared with TITERMAX adjuvant, from Vaxcell, Inc., and subsequent immunizations are prepared with incomplete Freund's adjuvant (IFA). At 11 weeks, the mice are IV boosted with 3-4 μ g VESPR in PBS. Three days after the IV boost, splenocytes are harvested and fused with an Ag8.653 myeloma fusion partner using 50% aqueous PEG 1500 solution. Hybridoma supernatants are screened for VESPR antibodies by dot blot assay against VESPR and an irrelevant Fc protein.

What is claimed is:

1. An isolated VESPR polypeptide that binds semaphorins.
2. A VESPR polypeptide comprising an amino acid sequence that is at least 90% identical to the amino acid sequence of SEQ ID NO:2, the VESPR polypeptide capable of binding semaphorins.
3. An isolated VESPR polypeptide of claim 2 wherein the semaphorin to which VESPR polypeptides is capable of binding is selected from the group consisting of A39R semaphorin and AHV semaphorin.
4. An isolated VESPR polypeptide encoded by DNA selected from the group consisting of:
 - (a) cDNA of SEQ ID NO:1; and
 - (b) DNA sequences that hybridize under moderately stringent conditions to the cDNA of (a); and which DNA sequences encode a polypeptide that binds semaphorins.
 - (c) DNA complementary to the DNA of (a) and (b).
5. A soluble VESPR polypeptide comprising an amino acid sequence selected from the group consisting of:
 - (a) amino acids x_1 to 945 of SEQ ID NO: 2, wherein x_1 is amino acid 1 or 35.
 - (b) a fragment of the sequence of (a), wherein the fragment is capable of binding a semaphorin.
6. The soluble VESPR polypeptide of claim 5 wherein the amino acid sequence is at least 90% identical to the sequence of (a) and the soluble VESPR polypeptide binds a semaphorin.
7. An isolated DNA encoding a VESPR polypeptide, the DNA selected from the group consisting of:
 - (a) DNA capable of hybridizing under highly stringent conditions to a nucleotide sequence consisting essentially of SEQ ID NO:1.

- (b) DNA capable of hybridizing under highly stringent conditions to DNA complementary to the sequence of (a),
 - (c) DNA sequences that, due to the degeneracy of the genetic code, is degenerate to the DNA of (a) and (b); and
 - (d) DNA complementary to the DNA of (a), (b), or (c).
8. An isolated DNA encoding a VESPR polypeptide, the DNA selected from the group consisting of:
- (a) DNA of SEQ ID NO:1;
 - (b) DNA sequences that hybridize under moderately stringent conditions to the cDNA of (a); and which DNA sequences encode a polypeptide that binds semaphorin;
 - (c) DNA sequences that, due to the degeneracy of the genetic code, encode VESPR polypeptides having the amino acid sequence of the polypeptides encoded by the DNA sequences of (a) or (b); and
 - (d) DNA complementary to the DNA of (a), (b), or (c).
9. An isolated DNA encoding an VESPR polypeptide wherein the VESPR polypeptide comprises an amino acid sequence that is at least 90% identical to the amino acid sequence of SEQ ID NO:2.
10. The isolated DNA of claim 9 wherein the VESPR polypeptide comprises the amino acid sequence of SEQ ID NO:2.
11. An isolated DNA encoding a soluble VESPR polypeptide, wherein the soluble VESPR polypeptide comprises an amino acid sequence that is at least 90% identical to a sequence selected from the group consisting of:
- (a) amino acids x_1 to 945 of SEQ ID NO:2, wherein x_1 is amino acid 1 or 35; and
 - (b) a fragment of the sequence of (a), wherein the soluble VESPR polypeptide binds a semaphorin.
12. A DNA of claim 11 wherein the soluble VESPR polypeptide comprises an amino acid sequence selected from the group consisting of:
- (a) amino acids x_1 to 945 of SEQ ID NO:2, wherein x_1 is amino acid 1 or 35; and
 - (b) a fragment of (a).

13. A fusion protein comprising amino acids x_1 to 945 where x_1 is amino acid 1 or 35 of SEQ ID NO:2
14. A recombinant expression vector comprising DNA of claim 7.
15. A process for preparing a VESPR polypeptide, the process comprising culturing a host cell transformed with an expression vector of claim 14 under conditions that promote expression of the polypeptide, and recovering the polypeptide.
16. A composition comprising a suitable diluent carrier and a polypeptide of claim 2.
17. An antibody that is immunoreactive with a polypeptide of claim 2.
18. A process for treating an inflammatory disease in a mammal afflicted with the disease, the process comprising administering an amount of VESPR polypeptide.
19. A method of separating cells having VESPR polypeptide on the surface thereof from a mixture of cells in suspension, comprising contacting the cells in the mixture with a contacting surface having a semaphorin, and separating the contacting surface and the suspension.

SEQUENCE LISTING

(1) GENERAL INFORMATION:

- (i) APPLICANT: Immunex Corporation, Melanie K. Spriggs,
Michael R. Comeau, Robert F. DuBose,
Richard S. Johnson
- (ii) TITLE OF INVENTION: VIRAL ENCODED SEMAPHORIN PROTEIN
RECEPTOR DNA AND POLYPEPTIDES
- (iii) NUMBER OF SEQUENCES: 10
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- (v) COMPUTER READABLE FORM:
 - (A) MEDIUM TYPE: Floppy disk
 - (B) COMPUTER: IBM PC compatible
 - (C) OPERATING SYSTEM: PC-DOS/MS-DOS
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(2) INFORMATION FOR SEQ ID NO:1:

- (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 4707 base pairs
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: double
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(iii) HYPOTHETICAL: NO

(iv) ANTI-SENSE: NO

(ix) FEATURE:

- (A) NAME/KEY: CDS
 (B) LOCATION: 1..4707

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:1:

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Ala Pro Leu Pro Leu Leu Ala Tyr Leu Leu Ala Leu Ala Ala Pro Gly	
20 25 30	
CGG GGC GCG GAC GAG CCC GTG TGG CGG TCG GAG CAA GCC ATC GGA GCC	144
Arg Gly Ala Asp Glu Pro Val Trp Arg Ser Glu Gln Ala Ile Gly Ala	
35 40 45	
ATC GCG GCG AGC CAG GAG GAC GGC GTG TTT GTG GCG AGC GGC AGC TGC	192
Ile Ala Ala Ser Gln Glu Asp Gly Val Phe Val Ala Ser Gly Ser Cys	
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CTG GAC CAG CTG GAC TAC AGC CTG GAG CAC AGC CTC TCG CGC CTG TAC	240
Leu Asp Gln Leu Asp Tyr Ser Leu Glu His Ser Leu Ser Arg Leu Tyr	
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Arg Cys Asn Pro Ala Ala Ser Asp His Asp Thr Ala Ile Ala Leu Lys	195	200	205	
GAC ACG GAG GGG CGC AGC CTG GCC ACG CAG GAG CTG GGG CGC CTC AAG				672
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Arg Ser Pro Thr Thr Thr Ala Leu Cys Leu Phe Arg Met Ser Glu Ile	325	330	335	
CAG GCG CGC GCC AAG AGG GTC AGC TGG GAC TTC AAG ACG GCC GAG AGC				1056
Gln Ala Arg Ala Lys Arg Val Ser Trp Asp Phe Lys Thr Ala Glu Ser	340	345	350	
CAC TGC AAA GAA GGG GAT CAA CCT GAA AGA GTC CAA CCA ATC GCA TCA				1104
His Cys Lys Glu Gly Asp Gln Pro Glu Arg Val Gln Pro Ile Ala Ser	355	360	365	
TCT ACC TTG ATC CAT TCC GAC CTG ACA TCC GTT TAT GGC ACC GTG GTA				1152
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Lys Val Ile Leu Gly Glu Asn Leu Thr Ser Asn Cys Pro Glu Val Ile	
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420 425 430	
CCT GTG AAG AAT ATC TAC ATT TAT CTA ACA GCT GGG AAA GAG GTG AGG	1344
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465 470 475 480	
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Cys Thr Phe Gln Gly Asp Cys Val His Ser Glu Asn Leu Glu Asn Trp	
485 490 495	
CTG GAT ATT TCG TCT GGA GCA AAA AAG TGC CCT AAA ATT CAG ATA ATT	1536
Leu Asp Ile Ser Ser Gly Ala Lys Lys Cys Pro Lys Ile Gln Ile Ile	
500 505 510	
CGA AGC AGT AAA GAA AAG ACT ACA GTG ACT ATG GTG GGA AGC TTC TCT	1584
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515 520 525	
CCA AGA CAC TCA AAG TGC ATG GTG AAG AAT GTG GAC TCT AGC AGG GAG	1632
Pro Arg His Ser Lys Cys Met Val Lys Asn Val Asp Ser Ser Arg Glu	
530 535 540	
CTC TGC CAG AAT AAA AGT CAG CCC AAC CGG ACC TGC ACC TGT AGC ATC	1680
Leu Cys Gln Asn Lys Ser Gln Pro Asn Arg Thr Cys Thr Cys Ser Ile	
545 550 555 560	
CCA ACC AGA GCA ACC TAC AAA GAT GTT TCA GTT GTC AAC GTG ATG TTC	1728
Pro Thr Arg Ala Thr Tyr Lys Asp Val Ser Val Val Asn Val Met Phe	
565 570 575	
TCC TTC GGT TCT TGG AAT TTA TCA GAC AGA TTC AAC TTT ACC AAC TGC	1776
Ser Phe Gly Ser Trp Asn Leu Ser Asp Arg Phe Asn Phe Thr Asn Cys	
580 585 590	
TCA TCA TTA AAA GAA TGC CCA GCA TGC GTA GAA ACT GGC TGC GCG TGG	1824
Ser Ser Leu Lys Glu Cys Pro Ala Cys Val Glu Thr Gly Cys Ala Trp	
595 600 605	
TGT AAA AGT GCA AGA AGG TGT ATC CAC CCC TTC ACA GCT TGC GAC CCT	1872
Cys Lys Ser Ala Arg Arg Cys Ile His Pro Phe Thr Ala Cys Asp Pro	

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TCT GAT TAT GAG AGA AAC CAG GAA CAG TGT CCA GTG GCT GTC GAG AAG Ser Asp Tyr Glu Arg Asn Gln Glu Gln Cys Pro Val Ala Val Glu Lys 625 630 635 640			1920
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TTT GAT GGT GGG AAC TGC TCT TCT GTG GGA TCC TTA TCC TAC ATT GCT Phe Asp Gly Gly Asn Cys Ser Ser Val Gly Ser Leu Ser Tyr Ile Ala 740 745 750			2256
CTG CCA CAT TGT TCC CTT ATA TTT CCT GCT ACC ACC TGG ATC AGT GGT Leu Pro His Cys Ser Leu Ile Phe Pro Ala Thr Thr Trp Ile Ser Gly 755 760 765			2304
GGT CAA AAT ATA ACC ATG ATG GGC AGA AAT TTT GAT GTA ATT GAC AAC Gly Gln Asn Ile Thr Met Met Gly Arg Asn Phe Asp Val Ile Asp Asn 770 775 780			2352
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AAA GAA AAT GAC AAC TTC AAT ATT TCC AAA AAA GAC ATT GAA ATT ACT Lys Glu Asn Asp Asn Phe Asn Ile Ser Lys Lys Asp Ile Glu Ile Thr 865 870 875 880	2640
CTC TTC CAT GGG GAA AAT GGG CAA TTA AAT TGC AGT TTT GAA AAT ATT Leu Phe His Gly Glu Asn Gly Gln Leu Asn Cys Ser Phe Glu Asn Ile 885 890 895	2688
ACT AGA AAT CAA GAT CTT ACC ACC ATC CTT TGC AAA ATT AAA GGC ATC Thr Arg Asn Gln Asp Leu Thr Thr Ile Leu Cys Lys Ile Lys Gly Ile 900 905 910	2736
AAG ACT GCA AGC ACC ATT GCC AAC TCT TCT AAG AAA GTT CGG GTC AAG Lys Thr Ala Ser Thr Ile Ala Asn Ser Ser Lys Lys Val Arg Val Lys 915 920 925	2784
CTG GGA AAC CTG GAG CTC TAC GTC GAG CAG GAG TCA GTT CCT TCC ACA Leu Gly Asn Leu Glu Leu Tyr Val Glu Gln Glu Ser Val Pro Ser Thr 930 935 940	2832
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GCG GCC GTG GGG GTG ACC AGG CAC AAA TCG AAG GAG CTG AGT CGC AAA Ala Ala Val Gly Val Thr Arg His Lys Ser Lys Glu Leu Ser Arg Lys 965 970 975	2928
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CGT GAC GGC TTT GCT GAG CTG CAG ATG GAT AAA TTG GAT GTG GTT GAT Arg Asp Gly Phe Ala Glu Leu Gln Met Asp Lys Leu Asp Val Val Asp 995 1000 1005	3024
AGT TTT GGA ACT GTT CCC TTC CTT GAC TAC AAA CAT TTT GCT CTG AGA Ser Phe Gly Thr Val Pro Phe Leu Asp Tyr Lys His Phe Ala Leu Arg 1010 1015 1020	3072
ACT TTC TTC CCT GAG TCA GGT GGC TTC ACC CAC ATC TTC ACT GAA GAT Thr Phe Phe Pro Glu Ser Gly Gly Phe Thr His Ile Phe Thr Glu Asp 1025 1030 1035 1040	3120
ATG CAT AAC AGA GAC GCC AAC GAC AAG AAT GAA AGT CTC ACA GCT TTG Met His Asn Arg Asp Ala Asn Asp Lys Asn Glu Ser Leu Thr Ala Leu 1045 1050 1055	3168
GAT GCC CTA ATC TGT AAT AAA AGC TTT CTT GTT ACT GTC ATC CAC ACC Asp Ala Leu Ile Cys Asn Lys Ser Phe Leu Val Thr Val Ile His Thr 1060 1065 1070	3216
CTT GAA AAG CAG AAG AAC TTT TCT GTG AAG GAC AGG TGT CTG TTT GCC Leu Glu Lys Gln Lys Asn Phe Ser Val Lys Asp Arg Cys Leu Phe Ala	3264

1075	1080	1085	
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ATC CTA GAG GTG CTG ACC AGG GAC TTG ATG GAA CAG TGT AGT AAC ATG Ile Leu Glu Val Leu Thr Arg Asp Leu Met Glu Gln Cys Ser Asn Met 1105 1110 1115 1120			3360
CAG CCG AAA CTC ATG CTG AGA CGC ACG GAG TCC GTC GTC GAA AAA CTC Gln Pro Lys Leu Met Leu Arg Arg Thr Glu Ser Val Val Glu Lys Leu 1125 1130 1135			3408
CTC ACA AAC TGG ATG TCC GTC TGC CTT TCT GGA TTT CTC CGG GAG ACT Leu Thr Asn Trp Met Ser Val Cys Leu Ser Gly Phe Leu Arg Glu Thr 1140 1145 1150			3456
GTC GGA GAG CCC TTC TAT TTG CTG GTG ACG ACT CTG AAC CAG AAA ATT Val Gly Glu Pro Phe Tyr Leu Leu Val Thr Thr Leu Asn Gln Lys Ile 1155 1160 1165			3504
AAC AAG GGT CCC GTG GAT GTA ATC ACT TGC AAA GCC CTG TAC ACA CTT Asn Lys Gly Pro Val Asp Val Ile Thr Cys Lys Ala Leu Tyr Thr Leu 1170 1175 1180			3552
AAT GAA GAC TGG CTG TTG TGG CAG GTT CCG GAA TTC AGT ACT GTG GCA Asn Glu Asp Trp Leu Leu Trp Gln Val Pro Glu Phe Ser Thr Val Ala 1185 1190 1195 1200			3600
TTA AAC GTC GTC TTT GAA AAA ATC CCG GAA AAC GAG AGT GCA GAT GTC Leu Asn Val Val Phe Glu Lys Ile Pro Glu Asn Glu Ser Ala Asp Val 1205 1210 1215			3648
TGT CGG AAT ATT TCA GTC AAT GTT CTC GAC TGT GAC ACC ATT GGC CAA Cys Arg Asn Ile Ser Val Asn Val Leu Asp Cys Asp Thr Ile Gly Gln 1220 1225 1230			3696
GCC AAA GAA AAG ATT TTC CAA GCA TTC TTA AGC AAA AAT GGC TCT CCT Ala Lys Glu Lys Ile Phe Gln Ala Phe Leu Ser Lys Asn Gly Ser Pro 1235 1240 1245			3744
TAT GGA CTT CAG CTT AAT GAA ATT GGT CTT GAG CTT CAA ATG GGC ACA Tyr Gly Leu Gln Leu Asn Glu Ile Gly Leu Glu Leu Gln Met Gly Thr 1250 1255 1260			3792
CGA CAG AAA GAA CTT CTG GAC ATC GAC AGT TCC TCC GTG ATT CTT GAA Arg Gln Lys Glu Leu Leu Asp Ile Asp Ser Ser Ser Val Ile Leu Glu 1265 1270 1275 1280			3840
GAT GGA ATC ACC AAG CTA AAC ACC ATT GGC CAC TAT GAG ATA TCA AAT Asp Gly Ile Thr Lys Leu Asn Thr Ile Gly His Tyr Glu Ile Ser Asn 1285 1290 1295			3888
GGA TCC ACT ATA AAA GTC TTT AAG AAG ATA GCA AAT TTT ACT TCA GAT Gly Ser Thr Ile Lys Val Phe Lys Lys Ile Ala Asn Phe Thr Ser Asp 1300 1305 1310			3936

GTG GAG TAC TCG GAT GAC CAC TGC CAT TTG ATT TTA CCA GAT TCG GAA Val Glu Tyr Ser Asp Asp His Cys His Leu Ile Leu Pro Asp Ser Glu 1315 1320 1325	3984
GCA TTC CAA GAT GTG CAA GGA AAG AGA CAT CGA GGG AAG CAC AAG TTC Ala Phe Gln Asp Val Gln Gly Lys Arg His Arg Gly Lys His Lys Phe 1330 1335 1340	4032
AAA GTA AAA GAA ATG TAT CTG ACA AAG CTG CTG TCG ACC AAG GTG GCA Lys Val Lys Glu Met Tyr Leu Thr Lys Leu Leu Ser Thr Lys Val Ala 1345 1350 1355 1360	4080
ATT CAT TCT GTG CTT GAA AAA CTT TTT AGA AGC ATT TGG AGT TTA CCC Ile His Ser Val Leu Glu Lys Leu Phe Arg Ser Ile Trp Ser Leu Pro 1365 1370 1375	4128
AAC AGC AGA GCT CCA TTT GCT ATA AAA TAC TTT TTT GAC TTT TTG GAC Asn Ser Arg Ala Pro Phe Ala Ile Lys Tyr Phe Phe Asp Phe Leu Asp 1380 1385 1390	4176
GCC CAG GCT GAA AAC AAA AAA ATC ACA GAT CCT GAC GTC GTA CAT ATT Ala Gln Ala Glu Asn Lys Lys Ile Thr Asp Pro Asp Val Val His Ile 1395 1400 1405	4224
TGG AAA ACA AAC AGC CTT CCT CTT CGC TTC TGG GTA AAC ATC CTG AAG Trp Lys Thr Asn Ser Leu Pro Leu Arg Phe Trp Val Asn Ile Leu Lys 1410 1415 1420	4272
AAC CCT CAG TTT GTC TTT GAC ATT AAG AAG ACA CCA CAT ATA GAC GGC Asn Pro Gln Phe Val Phe Asp Ile Lys Lys Thr Pro His Ile Asp Gly 1425 1430 1435 1440	4320
TGT TTG TCA GTG ATT GCC CAG GCA TTC ATG GAT GCA TTT TCT CTC ACA Cys Leu Ser Val Ile Ala Gln Ala Phe Met Asp Ala Phe Ser Leu Thr 1445 1450 1455	4368
GAG CAG CAA CTA GGG AAG GAA GCA CCA ACT AAT AAG CTT CTC TAT GCC Glu Gln Gln Leu Gly Lys Glu Ala Pro Thr Asn Lys Leu Leu Tyr Ala 1460 1465 1470	4416
AAG GAT ATC CCA ACC TAC AAA GAA GAA GTA AAA TCT TAT TAC AAA GCA Lys Asp Ile Pro Thr Tyr Lys Glu Glu Val Lys Ser Tyr Tyr Lys Ala 1475 1480 1485	4464
ATC AGG GAT TTG CCT CCA TTG TCA TCC TCA GAA ATG GAA GAA TTT TTA Ile Arg Asp Leu Pro Pro Leu Ser Ser Ser Glu Met Glu Glu Phe Leu 1490 1495 1500	4512
ACT CAG GAA TCT AAG AAA CAT GAA AAT GAA TTT AAT GAA GAA GTG GCC Thr Gln Glu Ser Lys Lys His Glu Asn Glu Phe Asn Glu Glu Val Ala 1505 1510 1515 1520	4560
TTG ACA GAA ATT TAC AAA TAC ATC GTA AAA TAT TTT GAT GAG ATT CTA Leu Thr Glu Ile Tyr Lys Tyr Ile Val Lys Tyr Phe Asp Glu Ile Leu 1525 1530 1535	4608
AAT AAA CTA GAA AGA GAA CGA GGG CTG GAA GAA GCT CAG AAA CAA CTC Asn Lys Leu Glu Arg Glu Arg Gly Leu Glu Glu Ala Gln Lys Gln Leu	4656

1540	1545	1550	
TTG CAT GTA AAA GTC TTA TTT GAT GAA AAG AAG AAA TGC AAG TGG ATG			4704
Leu His Val Lys Val Leu Phe Asp Glu Lys Lys Lys Cys Lys Trp Met			
1555	1560	1565	

TAA 4707
*

(2) INFORMATION FOR SEQ ID NO:2:

- (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 1569 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:2:

Met	Glu	Val	Ser	Arg	Arg	Lys	Ala	Pro	Pro	Arg	Pro	Pro	Arg	Pro	Ala
1				5				10						15	
Ala	Pro	Leu	Pro	Leu	Leu	Ala	Tyr	Leu	Leu	Ala	Leu	Ala	Ala	Pro	Gly
		20						25					30		
Arg	Gly	Ala	Asp	Glu	Pro	Val	Trp	Arg	Ser	Glu	Gln	Ala	Ile	Gly	Ala
		35					40						45		
Ile	Ala	Ala	Ser	Gln	Glu	Asp	Gly	Val	Phe	Val	Ala	Ser	Gly	Ser	Cys
	50					55					60				
Leu	Asp	Gln	Leu	Asp	Tyr	Ser	Leu	Glu	His	Ser	Leu	Ser	Arg	Leu	Tyr
	65				70					75				80	
Arg	Asp	Gln	Ala	Gly	Asn	Cys	Thr	Glu	Pro	Val	Ser	Leu	Ala	Pro	Pro
			85						90					95	
Ala	Arg	Pro	Arg	Pro	Gly	Ser	Ser	Phe	Ser	Lys	Leu	Leu	Leu	Pro	Tyr
		100						105					110		
Arg	Glu	Gly	Ala	Ala	Gly	Leu	Gly	Gly	Leu	Leu	Leu	Thr	Gly	Trp	Thr
		115				120						125			
Phe	Asp	Arg	Gly	Ala	Cys	Glu	Val	Arg	Pro	Leu	Gly	Asn	Leu	Ser	Arg
	130					135					140				
Asn	Ser	Leu	Arg	Asn	Gly	Thr	Glu	Val	Val	Ser	Cys	His	Pro	Gln	Gly
	145				150					155				160	
Ser	Thr	Ala	Gly	Val	Val	Tyr	Arg	Ala	Gly	Arg	Asn	Asn	Arg	Trp	Tyr
			165						170					175	
Leu	Ala	Val	Ala	Ala	Thr	Tyr	Val	Leu	Pro	Glu	Pro	Glu	Thr	Ala	Ser
		180						185					190		
Arg	Cys	Asn	Pro	Ala	Ala	Ser	Asp	His	Asp	Thr	Ala	Ile	Ala	Leu	Lys

195	200	205
Asp Thr Glu Gly Arg Ser Leu Ala Thr Gln Glu Leu Gly Arg Leu Lys 210 215 220		
Leu Cys Glu Gly Ala Gly Ser Leu His Phe Val Asp Ala Phe Leu Trp 225 230 235 240		
Asn Gly Ser Ile Tyr Phe Pro Tyr Tyr Pro Tyr Asn Tyr Thr Ser Gly 245 250 255		
Ala Ala Thr Gly Trp Pro Ser Met Ala Arg Ile Ala Gln Ser Thr Glu 260 265 270		
Val Leu Phe Gln Gly Gln Ala Ser Leu Asp Cys Gly His Gly His Pro 275 280 285		
Asp Gly Arg Arg Leu Leu Leu Ser Ser Ser Leu Val Glu Ala Leu Asp 290 295 300		
Val Trp Ala Gly Val Phe Ser Ala Ala Ala Gly Glu Gly Gln Glu Arg 305 310 315 320		
Arg Ser Pro Thr Thr Thr Ala Leu Cys Leu Phe Arg Met Ser Glu Ile 325 330 335		
Gln Ala Arg Ala Lys Arg Val Ser Trp Asp Phe Lys Thr Ala Glu Ser 340 345 350		
His Cys Lys Glu Gly Asp Gln Pro Glu Arg Val Gln Pro Ile Ala Ser 355 360 365		
Ser Thr Leu Ile His Ser Asp Leu Thr Ser Val Tyr Gly Thr Val Val 370 375 380		
Met Asn Arg Thr Val Leu Phe Leu Gly Thr Gly Asp Gly Gln Leu Leu 385 390 395 400		
Lys Val Ile Leu Gly Glu Asn Leu Thr Ser Asn Cys Pro Glu Val Ile 405 410 415		
Tyr Glu Ile Lys Glu Glu Thr Pro Val Phe Tyr Lys Leu Val Pro Asp 420 425 430		
Pro Val Lys Asn Ile Tyr Ile Tyr Leu Thr Ala Gly Lys Glu Val Arg 435 440 445		
Arg Ile Arg Val Ala Asn Cys Asn Lys His Lys Ser Cys Ser Glu Cys 450 455 460		
Leu Thr Ala Thr Asp Pro His Cys Gly Trp Cys His Ser Leu Gln Arg 465 470 475 480		
Cys Thr Phe Gln Gly Asp Cys Val His Ser Glu Asn Leu Glu Asn Trp 485 490 495		
Leu Asp Ile Ser Ser Gly Ala Lys Lys Cys Pro Lys Ile Gln Ile Ile 500 505 510		

Arg Ser Ser Lys Glu Lys Thr Thr Val Thr Met Val Gly Ser Phe Ser
 515 520 525
 Pro Arg His Ser Lys Cys Met Val Lys Asn Val Asp Ser Ser Arg Glu
 530 535 540
 Leu Cys Gln Asn Lys Ser Gln Pro Asn Arg Thr Cys Thr Cys Ser Ile
 545 550 555 560
 Pro Thr Arg Ala Thr Tyr Lys Asp Val Ser Val Val Asn Val Met Phe
 565 570 575
 Ser Phe Gly Ser Trp Asn Leu Ser Asp Arg Phe Asn Phe Thr Asn Cys
 580 585 590
 Ser Ser Leu Lys Glu Cys Pro Ala Cys Val Glu Thr Gly Cys Ala Trp
 595 600 605
 Cys Lys Ser Ala Arg Arg Cys Ile His Pro Phe Thr Ala Cys Asp Pro
 610 615 620
 Ser Asp Tyr Glu Arg Asn Gln Glu Gln Cys Pro Val Ala Val Glu Lys
 625 630 635 640
 Thr Ser Gly Gly Gly Arg Pro Lys Glu Asn Lys Gly Asn Arg Thr Asn
 645 650 655
 Gln Ala Leu Gln Val Phe Tyr Ile Lys Ser Ile Glu Pro Gln Lys Val
 660 665 670
 Ser Thr Leu Gly Lys Ser Asn Val Ile Val Thr Gly Ala Asn Phe Thr
 675 680 685
 Arg Ala Ser Asn Ile Thr Met Ile Leu Lys Gly Thr Ser Thr Cys Asp
 690 695 700
 Lys Asp Val Ile Gln Val Ser His Val Leu Asn Asp Thr His Met Lys
 705 710 715 720
 Phe Ser Leu Pro Ser Ser Arg Lys Glu Met Lys Asp Val Cys Ile Gln
 725 730 735
 Phe Asp Gly Gly Asn Cys Ser Ser Val Gly Ser Leu Ser Tyr Ile Ala
 740 745 750
 Leu Pro His Cys Ser Leu Ile Phe Pro Ala Thr Thr Trp Ile Ser Gly
 755 760 765
 Gly Gln Asn Ile Thr Met Met Gly Arg Asn Phe Asp Val Ile Asp Asn
 770 775 780
 Leu Ile Ile Ser His Glu Leu Lys Gly Asn Ile Asn Val Ser Glu Tyr
 785 790 795 800
 Cys Val Ala Thr Tyr Cys Gly Phe Leu Ala Pro Ser Leu Lys Ser Ser
 805 810 815

Lys Val Arg Thr Asn Val Thr Val Lys Leu Arg Val Gln Asp Thr Tyr
 820 825 830
 Leu Asp Cys Gly Thr Leu Gln Tyr Arg Glu Asp Pro Arg Phe Thr Gly
 835 840 845
 Tyr Arg Val Glu Ser Glu Val Asp Thr Glu Leu Glu Val Lys Ile Gln
 850 855 860
 Lys Glu Asn Asp Asn Phe Asn Ile Ser Lys Lys Asp Ile Glu Ile Thr
 865 870 875 880
 Leu Phe His Gly Glu Asn Gly Gln Leu Asn Cys Ser Phe Glu Asn Ile
 885 890 895
 Thr Arg Asn Gln Asp Leu Thr Thr Ile Leu Cys Lys Ile Lys Gly Ile
 900 905 910
 Lys Thr Ala Ser Thr Ile Ala Asn Ser Ser Lys Lys Val Arg Val Lys
 915 920 925
 Leu Gly Asn Leu Glu Leu Tyr Val Glu Gln Glu Ser Val Pro Ser Thr
 930 935 940
 Trp Tyr Phe Leu Ile Val Leu Pro Val Leu Leu Val Ile Val Ile Phe
 945 950 955 960
 Ala Ala Val Gly Val Thr Arg His Lys Ser Lys Glu Leu Ser Arg Lys
 965 970 975
 Gln Ser Gln Gln Leu Glu Leu Leu Glu Ser Glu Leu Arg Lys Glu Ile
 980 985 990
 Arg Asp Gly Phe Ala Glu Leu Gln Met Asp Lys Leu Asp Val Val Asp
 995 1000 1005
 Ser Phe Gly Thr Val Pro Phe Leu Asp Tyr Lys His Phe Ala Leu Arg
 1010 1015 1020
 Thr Phe Phe Pro Glu Ser Gly Gly Phe Thr His Ile Phe Thr Glu Asp
 1025 1030 1035 1040
 Met His Asn Arg Asp Ala Asn Asp Lys Asn Glu Ser Leu Thr Ala Leu
 1045 1050 1055
 Asp Ala Leu Ile Cys Asn Lys Ser Phe Leu Val Thr Val Ile His Thr
 1060 1065 1070
 Leu Glu Lys Gln Lys Asn Phe Ser Val Lys Asp Arg Cys Leu Phe Ala
 1075 1080 1085
 Ser Phe Leu Thr Ile Ala Leu Gln Thr Lys Leu Val Tyr Leu Thr Ser
 1090 1095 1100
 Ile Leu Glu Val Leu Thr Arg Asp Leu Met Glu Gln Cys Ser Asn Met
 1105 1110 1115 1120

Gln Pro Lys Leu Met Leu Arg Arg Thr Glu Ser Val Val Glu Lys Leu
 1125 1130 1135
 Leu Thr Asn Trp Met Ser Val Cys Leu Ser Gly Phe Leu Arg Glu Thr
 1140 1145 1150
 Val Gly Glu Pro Phe Tyr Leu Leu Val Thr Thr Leu Asn Gln Lys Ile
 1155 1160 1165
 Asn Lys Gly Pro Val Asp Val Ile Thr Cys Lys Ala Leu Tyr Thr Leu
 1170 1175 1180
 Asn Glu Asp Trp Leu Leu Trp Gln Val Pro Glu Phe Ser Thr Val Ala
 1185 1190 1195 1200
 Leu Asn Val Val Phe Glu Lys Ile Pro Glu Asn Glu Ser Ala Asp Val
 1205 1210 1215
 Cys Arg Asn Ile Ser Val Asn Val Leu Asp Cys Asp Thr Ile Gly Gln
 1220 1225 1230
 Ala Lys Glu Lys Ile Phe Gln Ala Phe Leu Ser Lys Asn Gly Ser Pro
 1235 1240 1245
 Tyr Gly Leu Gln Leu Asn Glu Ile Gly Leu Glu Leu Gln Met Gly Thr
 1250 1255 1260
 Arg Gln Lys Glu Leu Leu Asp Ile Asp Ser Ser Ser Val Ile Leu Glu
 1265 1270 1275 1280
 Asp Gly Ile Thr Lys Leu Asn Thr Ile Gly His Tyr Glu Ile Ser Asn
 1285 1290 1295
 Gly Ser Thr Ile Lys Val Phe Lys Lys Ile Ala Asn Phe Thr Ser Asp
 1300 1305 1310
 Val Glu Tyr Ser Asp Asp His Cys His Leu Ile Leu Pro Asp Ser Glu
 1315 1320 1325
 Ala Phe Gln Asp Val Gln Gly Lys Arg His Arg Gly Lys His Lys Phe
 1330 1335 1340
 Lys Val Lys Glu Met Tyr Leu Thr Lys Leu Leu Ser Thr Lys Val Ala
 1345 1350 1355 1360
 Ile His Ser Val Leu Glu Lys Leu Phe Arg Ser Ile Trp Ser Leu Pro
 1365 1370 1375
 Asn Ser Arg Ala Pro Phe Ala Ile Lys Tyr Phe Phe Asp Phe Leu Asp
 1380 1385 1390
 Ala Gln Ala Glu Asn Lys Lys Ile Thr Asp Pro Asp Val Val His Ile
 1395 1400 1405
 Trp Lys Thr Asn Ser Leu Pro Leu Arg Phe Trp Val Asn Ile Leu Lys
 1410 1415 1420

Asn Pro Gln Phe Val Phe Asp Ile Lys Lys Thr Pro His Ile Asp Gly	
1425	1440
Cys Leu Ser Val Ile Ala Gln Ala Phe Met Asp Ala Phe Ser Leu Thr	
1445	1455
Glu Gln Gln Leu Gly Lys Glu Ala Pro Thr Asn Lys Leu Leu Tyr Ala	
1460	1470
Lys Asp Ile Pro Thr Tyr Lys Glu Glu Val Lys Ser Tyr Tyr Lys Ala	
1475	1485
Ile Arg Asp Leu Pro Pro Leu Ser Ser Ser Glu Met Glu Glu Phe Leu	
1490	1500
Thr Gln Glu Ser Lys Lys His Glu Asn Glu Phe Asn Glu Glu Val Ala	
1505	1520
Leu Thr Glu Ile Tyr Lys Tyr Ile Val Lys Tyr Phe Asp Glu Ile Leu	
1525	1535
Asn Lys Leu Glu Arg Glu Arg Gly Leu Glu Glu Ala Gln Lys Gln Leu	
1540	1550
Leu His Val Lys Val Leu Phe Asp Glu Lys Lys Lys Cys Lys Trp Met	
1555	1565

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(2) INFORMATION FOR SEQ ID NO:3:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 31 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:3:

TGTCAGTAGT ATCGAATGGC ATAAGTTGA A

31

(2) INFORMATION FOR SEQ ID NO:4:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 33 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(iii) HYPOTHETICAL: NO

(iv) ANTI-SENSE: NO

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:4:

GACAGCGGCC GCCTATTACA TTTTAAGTAT TTT

33

(2) INFORMATION FOR SEQ ID NO:5:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 18 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: primer

(iii) HYPOTHETICAL: NO

(iv) ANTI-SENSE: NO

(ix) FEATURE:

- (A) NAME/KEY:
- (B) LOCATION:

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:5:

GCGGGACTCA GAGTCACC

18

(2) INFORMATION FOR SEQ ID NO:6:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 43 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: primer

(iii) HYPOTHETICAL: NO

(iv) ANTI-SENSE: NO

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:6:

GGATCCTAAT ACGACTCACT ATAGGGAGGA AACCACTCCG AAC

43

(2) INFORMATION FOR SEQ ID NO:7:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 1983 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(iii) HYPOTHETICAL: NO

(iv) ANTI-SENSE: NO

(ix) FEATURE:

(A) NAME/KEY: CDS

(B) LOCATION: 1..1983

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:7:

ATG TTC CAT GTT TCT TTT AGA TAT ATC TTT GGA ATT CCT CCA CTG ATC	48
Met Phe His Val Ser Phe Arg Tyr Ile Phe Gly Ile Pro Pro Leu Ile	
1 5 10 15	
CTT GTT CTG CTG CCT GTC ACT AGC TCT GAC TAC AAA GAT GAC GAT GAT	96
Leu Val Leu Leu Pro Val Thr Ser Ser Asp Tyr Lys Asp Asp Asp Asp	
20 25 30	
AAA AGA TCT TGT GAC AAA ACT CAC ACA TGC CCA CCG TGC CCA GCA CCT	144
Lys Arg Ser Cys Asp Lys Thr His Thr Cys Pro Pro Cys Pro Ala Pro	
35 40 45	
GAA GCC GAG GGC GCG CCG TCA GTC TTC CTC TTC CCC CCA AAA CCC AAG	192
Glu Ala Glu Gly Ala Pro Ser Val Phe Leu Phe Pro Pro Lys Pro Lys	
50 55 60	
GAC ACC CTC ATG ATC TCC CGG ACC CCT GAG GTC ACA TGC GTG GTG GTG	240
Asp Thr Leu Met Ile Ser Arg Thr Pro Glu Val Thr Cys Val Val Val	
65 70 75 80	
GAC GTG AGC CAC GAA GAC CCT GAG GTC AAG TTC AAC TGG TAC GTG GAC	288
Asp Val Ser His Glu Asp Pro Glu Val Lys Phe Asn Trp Tyr Val Asp	
85 90 95	
GGC GTG GAG GTG CAT AAT GCC AAG ACA AAG CCG CGG GAG GAG CAG TAC	336
Gly Val Glu Val His Asn Ala Lys Thr Lys Pro Arg Glu Glu Gln Tyr	
100 105 110	
AAC AGC ACG TAC CGT GTG GTC AGC GTC CTC ACC GTC CTG CAC CAG GAC	384
Asn Ser Thr Tyr Arg Val Val Ser Val Leu Thr Val Leu His Gln Asp	
115 120 125	
TGG CTG AAT GGC AAG GAG TAC AAG TGC AAG GTC TCC AAC AAA GCC CTC	432
Trp Leu Asn Gly Lys Glu Tyr Lys Cys Lys Val Ser Asn Lys Ala Leu	
130 135 140	
CCA GCC CCC ATC GAG AAA ACC ATC TCC AAA GCC AAA GGG CAG CCC CGA	480
Pro Ala Pro Ile Glu Lys Thr Ile Ser Lys Ala Lys Gly Gln Pro Arg	
145 150 155 160	
GAA CCA CAG GTG TAC ACC CTG CCC CCA TCC CGG GAG GAG ATG ACC AAG	528
Glu Pro Gln Val Tyr Thr Leu Pro Pro Ser Arg Glu Glu Met Thr Lys	
165 170 175	

AAC CAG GTC AGC CTG ACC TGC CTG GTC AAA GGC TTC TAT CCC AGC GAC Asn Gln Val Ser Leu Thr Cys Leu Val Lys Gly Phe Tyr Pro Ser Asp 180 185 190	576
ATC GCC GTG GAG TGG GAG AGC AAT GGG CAG CCG GAG AAC AAC TAC AAG Ile Ala Val Glu Trp Glu Ser Asn Gly Gln Pro Glu Asn Asn Tyr Lys 195 200 205	624
ACC ACG CCT CCC GTG CTG GAC TCC GAC GGC TCC TTC TTC CTC TAT AGC Thr Thr Pro Pro Val Leu Asp Ser Asp Gly Ser Phe Phe Leu Tyr Ser 210 215 220	672
AAG CTC ACC GTG GAC AAG AGC AGG TGG CAG CAG GGG AAC GTC TTC TCA Lys Leu Thr Val Asp Lys Ser Arg Trp Gln Gln Gly Asn Val Phe Ser 225 230 235 240	720
TGC TCC GTG ATG CAT GAG GCT CTG CAC AAC CAC TAC ACG CAG AAG AGC Cys Ser Val Met His Glu Ala Leu His Asn His Tyr Thr Gln Lys Ser 245 250 255	768
CTC TCC CTG TCT CCG GGT AAA GGA GGG GGC GGA TCA GGG GGC GGA GGA Leu Ser Leu Ser Pro Gly Lys Gly Gly Gly Gly Ser Gly Gly Gly Gly 260 265 270	816
TCT ACT AGT ATC GAA TGG CAT AAG TTT GAA ACG AGT GAA GAA ATA ATT Ser Thr Ser Ile Glu Trp His Lys Phe Glu Thr Ser Glu Glu Ile Ile 275 280 285	864
TCT ACT TAC TTA ATA GAT GAT GTA TTA TAC ACG GGC GTT AAT GGG GCG Ser Thr Tyr Leu Ile Asp Asp Val Leu Tyr Thr Gly Val Asn Gly Ala 290 295 300	912
GTA TAT ACA TTT TCA AAT AAT GAA CTA AAC AAA ACT GGT TTA ACT AAT Val Tyr Thr Phe Ser Asn Asn Glu Leu Asn Lys Thr Gly Leu Thr Asn 305 310 315 320	960
AAC AAT AAT TAT ATC ACA ACA TCT ATA AAA GTA GAG GAT ACA TTA GTA Asn Asn Asn Tyr Ile Thr Thr Ser Ile Lys Val Glu Asp Thr Leu Val 325 330 335	1008
TGC GGA ACC AAT AAC GGA AAC CCC AAA TGT TGG AAA ATA GAC GGT TCC Cys Gly Thr Asn Asn Gly Asn Pro Lys Cys Trp Lys Ile Asp Gly Ser 340 345 350	1056
GAA GAT CCA AAA TAT AGA GGT AGA GGA TAT GCT CCT TAT CAA AAT AGT Glu Asp Pro Lys Tyr Arg Gly Arg Gly Tyr Ala Pro Tyr Gln Asn Ser 355 360 365	1104
AAA GTG ACG ATA ATC AGT CAT AAC GAA TGT GTA CTA TCT GAT ATA AAC Lys Val Thr Ile Ile Ser His Asn Glu Cys Val Leu Ser Asp Ile Asn 370 375 380	1152
ATA TCA AAA GAA GGA ATT AAA AGA TGG AGA AGA TTT GAC GGA CCA TGT Ile Ser Lys Glu Gly Ile Lys Arg Trp Arg Arg Phe Asp Gly Pro Cys 385 390 395 400	1200

GGT TAT GAT TTA TAC ACG GCA GAT AAC GTG ATT CCA AAA GAT GGT GTG	1248
Gly Tyr Asp Leu Tyr Thr Ala Asp Asn Val Ile Pro Lys Asp Gly Val	
405 410 415	
CGT GGA GCA TTC GTT GAT AAA GAC GGC ACT TAT GAC AAA GTT TAC ATT	1296
Arg Gly Ala Phe Val Asp Lys Asp Gly Thr Tyr Asp Lys Val Tyr Ile	
420 425 430	
CTT TTC ACT GAT ACT ATC GAC ACA AAG AGA ATT GTT AAA ATT CCG TAT	1344
Leu Phe Thr Asp Thr Ile Asp Thr Lys Arg Ile Val Lys Ile Pro Tyr	
435 440 445	
ATA GCA CAA ATG TGC TTA AAT GAC GAA GGT GGT CCA TCA TCA TTG TCT	1392
Ile Ala Gln Met Cys Leu Asn Asp Glu Gly Gly Pro Ser Ser Leu Ser	
450 455 460	
AGT CAT AGA TGG TCG ACG TTT CTC AAG GTC GAA TTA GAA TGT GAT ATC	1440
Ser His Arg Trp Ser Thr Phe Leu Lys Val Glu Leu Glu Cys Asp Ile	
465 470 475 480	
GAC GGA AGA AGT TAT AGA CAA ATT ATT CAT TCT AAA GCT ATA AAA ACA	1488
Asp Gly Arg Ser Tyr Arg Gln Ile Ile His Ser Lys Ala Ile Lys Thr	
485 490 495	
GAT AAT GAT ACG ATA CTA TAT GTA TTC TTT GAT AGT CCT TAT TCC AAG	1536
Asp Asn Asp Thr Ile Leu Tyr Val Phe Phe Asp Ser Pro Tyr Ser Lys	
500 505 510	
TCC GCA TTA TGT ACC TAT TCT ATG AAT GCC ATT AAA CAC TCT TTT TCT	1584
Ser Ala Leu Cys Thr Tyr Ser Met Asn Ala Ile Lys His Ser Phe Ser	
515 520 525	
ACG TCA AAA TTG GGA GGA TAT ACA AAG CAA TTG CCG TCT CCA GCT CCT	1632
Thr Ser Lys Leu Gly Gly Tyr Thr Lys Gln Leu Pro Ser Pro Ala Pro	
530 535 540	
GGT ATA TGT CTA CCA GCT GGA AAA GTT GTT CCA CAT ACC ACG TTT GAC	1680
Gly Ile Cys Leu Pro Ala Gly Lys Val Val Pro His Thr Thr Phe Asp	
545 550 555 560	
ATC ATA GAA CAA TAT AAT GAG CTA GAT GAT ATT ATA AAG CCT TTA TCT	1728
Ile Ile Glu Gln Tyr Asn Glu Leu Asp Asp Ile Ile Lys Pro Leu Ser	
565 570 575	
CAA CCT ATC TTC GAA GGA CCG TCT GGT GTT AAA TGG TTC GAT ATA AAG	1776
Gln Pro Ile Phe Glu Gly Pro Ser Gly Val Lys Trp Phe Asp Ile Lys	
580 585 590	
GAG AAG GAA AAT GAA CAT CGG GAA TAT AGA ATA TAC TTC ATA AAA GAA	1824
Glu Lys Glu Asn Glu His Arg Glu Tyr Arg Ile Tyr Phe Ile Lys Glu	
595 600 605	
AAT ACT ATA TAT TCG TTC GAT ACA AAA TCT AAA CAA ACT CGT AGT GCA	1872
Asn Thr Ile Tyr Ser Phe Asp Thr Lys Ser Lys Gln Thr Arg Ser Ala	
610 615 620	

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CAA GTT GAT GCG CGA CTA TTT TCA GTA ATG GTA ACT TCG AAA CCG TTA 1920
Gln Val Asp Ala Arg Leu Phe Ser Val Met Val Thr Ser Lys Pro Leu
625                      630                      635                      640

TTT ATA GCA GAT ATA GGG ATA GGA GTA GGA ATA CCA CGA ATG AAA AAA 1968
Phe Ile Ala Asp Ile Gly Ile Gly Val Gly Ile Pro Arg Met Lys Lys
645                      650                      655

ATA CTT AAA ATG TAA
Ile Leu Lys Met *
660
1983

```

(2) INFORMATION FOR SEQ ID NO:8:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 661 amino acids
- (B) TYPE: amino acid
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:8:

```

Met Phe His Val Ser Phe Arg Tyr Ile Phe Gly Ile Pro Pro Leu Ile
 1              5              10              15

Leu Val Leu Leu Pro Val Thr Ser Ser Asp Tyr Lys Asp Asp Asp Asp
20              25              30

Lys Arg Ser Cys Asp Lys Thr His Thr Cys Pro Pro Cys Pro Ala Pro
35              40              45

Glu Ala Glu Gly Ala Pro Ser Val Phe Leu Phe Pro Pro Lys Pro Lys
50              55              60

Asp Thr Leu Met Ile Ser Arg Thr Pro Glu Val Thr Cys Val Val Val
65              70              75              80

Asp Val Ser His Glu Asp Pro Glu Val Lys Phe Asn Trp Tyr Val Asp
85              90              95

Gly Val Glu Val His Asn Ala Lys Thr Lys Pro Arg Glu Glu Gln Tyr
100             105             110

Asn Ser Thr Tyr Arg Val Val Ser Val Leu Thr Val Leu His Gln Asp
115             120             125

Trp Leu Asn Gly Lys Glu Tyr Lys Cys Lys Val Ser Asn Lys Ala Leu
130             135             140

Pro Ala Pro Ile Glu Lys Thr Ile Ser Lys Ala Lys Gly Gln Pro Arg
145             150             155             160

Glu Pro Gln Val Tyr Thr Leu Pro Pro Ser Arg Glu Glu Met Thr Lys
165             170             175

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Asn Gln Val Ser Leu Thr Cys Leu Val Lys Gly Phe Tyr Pro Ser Asp
 180 185 190
 Ile Ala Val Glu Trp Glu Ser Asn Gly Gln Pro Glu Asn Asn Tyr Lys
 195 200 205
 Thr Thr Pro Pro Val Leu Asp Ser Asp Gly Ser Phe Phe Leu Tyr Ser
 210 215 220
 Lys Leu Thr Val Asp Lys Ser Arg Trp Gln Gln Gly Asn Val Phe Ser
 225 230 235 240
 Cys Ser Val Met His Glu Ala Leu His Asn His Tyr Thr Gln Lys Ser
 245 250 255
 Leu Ser Leu Ser Pro Gly Lys Gly Gly Gly Ser Gly Gly Gly Gly
 260 265 270
 Ser Thr Ser Ile Glu Trp His Lys Phe Glu Thr Ser Glu Glu Ile Ile
 275 280 285
 Ser Thr Tyr Leu Ile Asp Asp Val Leu Tyr Thr Gly Val Asn Gly Ala
 290 295 300
 Val Tyr Thr Phe Ser Asn Asn Glu Leu Asn Lys Thr Gly Leu Thr Asn
 305 310 315 320
 Asn Asn Asn Tyr Ile Thr Thr Ser Ile Lys Val Glu Asp Thr Leu Val
 325 330 335
 Cys Gly Thr Asn Asn Gly Asn Pro Lys Cys Trp Lys Ile Asp Gly Ser
 340 345 350
 Glu Asp Pro Lys Tyr Arg Gly Arg Gly Tyr Ala Pro Tyr Gln Asn Ser
 355 360 365
 Lys Val Thr Ile Ile Ser His Asn Glu Cys Val Leu Ser Asp Ile Asn
 370 375 380
 Ile Ser Lys Glu Gly Ile Lys Arg Trp Arg Arg Phe Asp Gly Pro Cys
 385 390 395 400
 Gly Tyr Asp Leu Tyr Thr Ala Asp Asn Val Ile Pro Lys Asp Gly Val
 405 410 415
 Arg Gly Ala Phe Val Asp Lys Asp Gly Thr Tyr Asp Lys Val Tyr Ile
 420 425 430
 Leu Phe Thr Asp Thr Ile Asp Thr Lys Arg Ile Val Lys Ile Pro Tyr
 435 440 445
 Ile Ala Gln Met Cys Leu Asn Asp Glu Gly Gly Pro Ser Ser Leu Ser
 450 455 460
 Ser His Arg Trp Ser Thr Phe Leu Lys Val Glu Leu Glu Cys Asp Ile
 465 470 475 480

Asp Gly Arg Ser Tyr Arg Gln Ile Ile His Ser Lys Ala Ile Lys Thr
 485 490 495
 Asp Asn Asp Thr Ile Leu Tyr Val Phe Phe Asp Ser Pro Tyr Ser Lys
 500 505 510
 Ser Ala Leu Cys Thr Tyr Ser Met Asn Ala Ile Lys His Ser Phe Ser
 515 520 525
 Thr Ser Lys Leu Gly Gly Tyr Thr Lys Gln Leu Pro Ser Pro Ala Pro
 530 535 540
 Gly Ile Cys Leu Pro Ala Gly Lys Val Val Pro His Thr Thr Phe Asp
 545 550 555 560
 Ile Ile Glu Gln Tyr Asn Glu Leu Asp Asp Ile Ile Lys Pro Leu Ser
 565 570 575
 Gln Pro Ile Phe Glu Gly Pro Ser Gly Val Lys Trp Phe Asp Ile Lys
 580 585 590
 Glu Lys Glu Asn Glu His Arg Glu Tyr Arg Ile Tyr Phe Ile Lys Glu
 595 600 605
 Asn Thr Ile Tyr Ser Phe Asp Thr Lys Ser Lys Gln Thr Arg Ser Ala
 610 615 620
 Gln Val Asp Ala Arg Leu Phe Ser Val Met Val Thr Ser Lys Pro Leu
 625 630 635 640
 Phe Ile Ala Asp Ile Gly Ile Gly Val Gly Ile Pro Arg Met Lys Lys
 645 650 655
 Ile Leu Lys Met *
 660

(2) INFORMATION FOR SEQ ID NO:9:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 33 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: primer

(iii) HYPOTHETICAL: NO

(iv) ANTI-SENSE: NO

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:9:

ATCGCATCAT CTACCTTCAT CCATTCCGAC CTG

33

(2) INFORMATION FOR SEQ ID NO:10:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 33 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: primer

(iii) HYPOTHETICAL: NO

(iv) ANTI-SENSE: NO

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:10:

TAAACACTCC GAACAGGATT TATGTTTATT GCA

33

INTERNATIONAL SEARCH REPORT

International Application No PCT/US 98/22879		
A. CLASSIFICATION OF SUBJECT MATTER IPC 6 C12N15/12 C07K14/705 C12N15/62 C07K16/28 C12P21/02 A61K38/17		
According to International Patent Classification (IPC) or to both national classification and IPC		
B. FIELDS SEARCHED Minimum documentation searched (classification system followed by classification symbols) IPC 6 C12N C07K C12P A61K		
Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched		
Electronic data base consulted during the international search (name of data base and, where practical, search terms used)		
C. DOCUMENTS CONSIDERED TO BE RELEVANT		
Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
P,X	COMEAU M R ET AL: "A poxvirus-encoded semaphorin induces cytokine production from monocytes and binds to a novel cellular semaphorin receptor, VESPR." IMMUNITY, (1998 APR) 8 (4) 473-82. JOURNAL CODE: CCF. ISSN: 1074-7613., XP002094205 United States see the whole document ---	1-17,19
X	ADAMS MD ET AL: "Human brain expressed sequence Tag EST01773." GENESEQ Database entry Q61322 Accession number Q61322; 16 Mar 1994 XP002094206 see sequence	7,8
X	& WO 93 16178 A (US DEPT HEALTH & HUMAN SERVICE) 19 August 1993 see page 473; example 4 ---	7,8
-/--		
<div style="display: flex; justify-content: space-between;"> <input checked="" type="checkbox"/> Further documents are listed in the continuation of box C. <input checked="" type="checkbox"/> Patent family members are listed in annex. </div>		
* Special categories of cited documents :		
<div style="display: flex; justify-content: space-between;"> <div style="width: 45%;"> <p>"A" document defining the general state of the art which is not considered to be of particular relevance</p> <p>"E" earlier document but published on or after the international filing date</p> <p>"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)</p> <p>"O" document referring to an oral disclosure, use, exhibition or other means</p> <p>"P" document published prior to the international filing date but later than the priority date claimed</p> </div> <div style="width: 45%;"> <p>"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention</p> <p>"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone</p> <p>"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.</p> <p>"&" document member of the same patent family</p> </div> </div>		
Date of the actual completion of the international search <div style="text-align: center; font-weight: bold;">22 February 1999</div>		Date of mailing of the international search report <div style="text-align: center; font-weight: bold;">09/03/1999</div>
Name and mailing address of the ISA European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Tx. 31 651 epo nl. Fax: (+31-70) 340-3016		Authorized officer <div style="text-align: center; font-weight: bold;">Esen, J</div>

INTERNATIONAL SEARCH REPORT

Internatic Application No
PCT/US 98/22879

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
Category :	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	<p>WO 95 07706 A (UNIV CALIFORNIA) 23 March 1995</p> <p>-----</p>	

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US 98/22879

Box I Observations where certain claims were found unsearchable (Continuation of Item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☒ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:
Remark: Although claim 18
is directed to a method of treatment of the human/animal
body, the search has been carried out and based on the alleged
effects of the compound/composition.
2. ☐ Claims Nos.:
because they relate to parts of the International Application that do not comply with the prescribed requirements to such
an extent that no meaningful International Search can be carried out, specifically:
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of Item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this International Search Report covers all
searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment
of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this International Search Report
covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is
restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
- ☐ No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No

PCT/US 98/22879

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 9507706 A	23-03-1995	US 5639856 A	17-06-1997
		AU 683494 B	13-11-1997
		AU 7724094 A	03-04-1995
		CA 2171638 A	23-03-1995
		EP 0721342 A	17-07-1996
		JP 9505725 T	10-06-1997
		US 5807826 A	15-09-1998
<hr/>			